

EMULSION CHAMBER TECHNOLOGY EXPERIMENT (ECT)

Final Report on Contract NAS8-38428

SUMMARY

The experimental objective of ECT was to develop space-borne emulsion chamber technology so that cosmic rays and nuclear interactions may subsequently be studied at extremely high energies with long exposures in space.

A small emulsion chamber was built and flown on flight STS-62 of the *Columbia* in March 1994. Analysis of the several hundred layers of radiation-sensitive material has shown excellent post-flight condition and suitability for cosmic ray physics analysis at much longer exposures. Temperature control of the stack was $20 \pm 1^\circ\text{C}$ throughout the active control period and no significant deviations of temperature or pressure in the chamber were observed over the entire mission operations period. The unfortunate flight attitude of the orbiter (almost 90% Earth viewing) prevented any significant number of heavy particles ($Z \geq 10$) reaching the stack and the inverted flow of shower particles in the calorimeter has not allowed evaluation of absolute primary cosmic ray-detection efficiency nor of the practical time limits of useful exposure of these calorimeters in space to the level of detail originally planned. Nevertheless, analysis of the observed backgrounds and quality of the processed photographic and plastic materials after the flight show that productive exposures of emulsion chambers are feasible in low orbit for periods of up to one year or longer. The engineering approaches taken in the ECT program were proven effective and no major environmental obstacles to prolonged flight are evident.

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1. Introduction

Emulsion chambers have proven an efficient means of measuring the charge composition and energy spectrum of cosmic rays in the region above 10^{13} eV (ref 1,2,3,4)). Such measurements require detector exposure factors of thousands of m^2 hours exposure above the atmosphere to provide adequate statistics. So far these have been, and continue to be, obtained using balloons. However, definitive measurements of abundances above 10^{15} eV require exposures of detectors of area several m^2 for periods of many months, and ultimately will require flight on an orbital platform. An engineering test flight of a $40 \times 50 \text{ cm}^2$ emulsion chamber was undertaken on the Space Shuttle to evaluate the effects of radiation background, launch, thermal and other environments on a heavy calorimeter of this type. Since the detector stack was composed of parallel sheets of total mass-thickness $\sim 120 \text{ g cm}^{-2}$, it also provided a thick structure easily modeled for radiation transport calculations. The stack included arrays of small dosimetry detectors to provide a means of calibrating the transport codes in the orbital radiation field at high shielding depths.

This paper describes the technical results of the first orbital flight of an emulsion chamber. The flight experiment was performed in March, 1994, on NASA's Space Shuttle *Columbia*, and designated STS-62. The Emulsion Chamber Technology (ECT) mission was planned to *assess the major uncertainties of space exposure of an emulsion calorimeter* through the flight of one sub-unit of a chamber. Deployment of an actual experiment for astrophysical research purposes would require an assembly of many such chambers. Verification of the effectiveness of the emulsion chamber technology in space is vital for large-scale cosmic ray experiments on the STS and the International Space Station platforms. The secondary objective of the ECT flight was to obtain radiation data for studying the effects of shielding on the penetrative particles of interest, utilizing the emulsion chamber's self-shielding materials that are much greater than the maximum value in ordinary manned space vehicles.

Our emulsion chambers contain a fine-grain, three-dimensional tracking telescope and a sampling calorimeter, which have proven powerful in observing very high energy cosmic ray protons and nuclei. This method was well established for balloon flight experiments by the Japanese American Cooperative Emulsion Experiment (JACEE) collaboration for direct observation of cosmic rays toward the "knee" region ($E \sim 10^{15}$ eV) (1). Very low intensities of cosmic rays in the "knee" region ($\sim 100/\text{m}^2 \text{ year sr}$ at $E \sim 10^{15}$ eV) call for a large-area and long-duration space experiments.

Long-duration space experiments, however, pose several logistic and technological challenges for emulsion chambers. Among many issues, the following three must at least be studied and overcome for successful experiments: (1) assess the feasibility of track registration and analyses with emulsions and X-ray films under conditions of high background density of slow protons coming from the radiation belts, (2) protection from hostile thermal environments to secure uniform and regular track registration quality, and (3) provision of the mechanical strength for safeguard of emulsions from extreme vibration and shock loads during launch of the Shuttle.

2. Instrumentation

The ECT emulsion chamber utilized over 120 double-sided emulsion plates with 70 X-ray films, 20 sheets of CR-39 plate, and 12 radiation lengths of lead absorbers (Fig. 1). A complete description of the flight stack configuration is given in Appendix D. The vertical material

thickness was 120 g/cm². All the materials were tightly mounted in a hermetically sealed, anodized aluminum chamber which was milled out of 2 blocks (Fig. 2). The ceiling or lid was an aluminum honeycomb panel. Thin Kapton-film heaters were mounted on the top and the bottom of the chamber. A dozen thermistors were mounted inside and outside the chamber, which, with an electronic control unit, regulated the uniform temperature of $20.0 \pm 0.1^{\circ}\text{C}$, at all the points in the chamber (Fig. 3) during the period while the system was powered. Ground procedures were designed so that the chamber temperature should not exceed 30°C at any time. No excursions above 24°C were encountered.

ECT was mounted in the *Columbia* on a cross-bay structure termed the Mission Particular Experiment Support Structure (MPESS) which was configured by Goddard Space Flight Center to carry a total of six experiments for the Office of Aeronautics and Space Technology (OAST). This configuration was termed OAST-2. Figs. 4, 5, and 6 show ECT and the OAST-2 in the bay of *Columbia*. The other experiments are identified in Fig. 7.

3. Flight Profile of STS-62

STS-62 was launched at 7:53 a.m. CST on March 4, 1994. The mission lasted a total of 335.3 hours or 13.97 days. The orbiter *Columbia* had a mean altitude of 296 km and an orbital inclination of 39.0 deg. While the cargo bay doors were open for most of the flight, the bay was mostly facing the Earth.

ECT was designed to measure cosmic rays and should ideally always have been facing deep space and away from Earth. Practically, on the Space Shuttle, that is never possible for a variety of reasons. At the time of mission planning, OAST-2 was secondary payload consisting of 6 experiments. The original ECT requirement was 80 hours space viewing, based on 50% Space:50% Earth viewing over a possible 6 to 7 day flight. This minimum was subsequently reduced to 35-40 hours deep-space viewing. Later, the mission was extended to 14 days without changing the ECT minimums. Because of United States Microgravity Payload (USMP) requirements and thermal problems with at least two OAST-2 payloads, the mission was changed from a basically gravity-gradient (-XLV) orientation to a mostly Earth-viewing flight. More than 10 days were spent in this latter orientation (-ZLV), with about 80 hours -XLV (equivalent to 40 hours deep-space). There were additional periods of several hours uninterrupted deep-space viewing which permitted testing of the cold case thermal control system but did not substantially add to the deep-space viewing fraction of the total exposure.

The result of this orientation mix was that only 12% of the orbital time was spent facing deep space, with the consequence that almost 90% of all cosmic ray showers entered ECT from the bottom, with most of the heavies having already interacted with materials in the Shuttle or the ECT support structure.

This produced two major impacts:

1. The fraction of heavy nuclei detected was very low compared with that originally expected.
2. The effect of retrograde showers through the calorimeter was quite different from normal (i.e. balloon) experience, preventing direct comparison of proton and helium fluxes and thus calibration of event-retrieval efficiency. Also, the particle

background distribution is different in the stack making extrapolations to much longer flight times of several months problematical.

The complex mass distribution of the materials in the Shuttle bay (see Figs. 8 and 9) has prevented the planned simplification of calculations using NASA radiation-transport codes.

It should be emphasized however that, despite these complications to our original analysis plan, many events were traced, and detailed analyses performed as discussed below. The engineering approaches used in the ECT experiment were adequate to protect the emulsion materials and would provide the design basis for a space exposure using emulsion calorimetry for a flight period many times longer.

4. Data Analysis

4.a) Materials for Data Analysis

All the photographic plates and solid state track detectors in the ECT were analyzed. To evaluate the performance of the emulsion chamber materials in space flight, an approximately-identical emulsion chamber of the 1994 Antarctic balloon flight experiment (JACEE-12: a 10-day circumpolar flight) performed only 3 months prior to the STS-62 flight, was analyzed for comparison.

The differences observed in these materials are largely a result of the thermal, mechanical and radiation environments of the space flight and a balloon flight. Table A1 illustrates the major differences of the ECT shuttle flight and the Antarctic balloon flight exposure. The Antarctic balloon flight was made in the stratosphere (~38 km above the ground) and did not receive any trapped-belt proton background. However, the year 1994, in which both STS-62 and JACEE-12 were flown, was close to solar minimum and background particle fluxes were high in both ECT and JACEE-12 chambers, due to trapped particles and low energy cosmic rays respectively. Comparisons of materials from these two flight materials exposed at solar-minimum help define the background problem and give solid bases for projecting the emulsion chamber capabilities for future long-duration orbital flights.

Materials in the Ground Control Unit for the ECT's STS-62 flight were developed and analyzed together with the flight materials. Other materials used in the analysis include those used in the Materials Compatibility Tests that were performed during the Production Phase (1991 - 93).

Table A1. Flight parameters of the STS-62 (ECT) and the Antarctic circumpolar experiment (JACEE-12).

	Flight Dates	Duration of Flight	Altitude (Average)	Temperature during Flight	Developments Processed in
ECT	3/4~18/94	14 Days	296 km	20 ± 0.1 °C	May, 1994
JACEE-12	1/3~14/94	10 Days	38 km	-5 ± 3 °C*	April, 1994

*Diurnal temperature variation at a particular plate. This does not include stable variations in plate temperatures from top to bottom (~15°C) in the stack, nor the cool-down period (~1 day) immediately following balloon launch.

4.b) X-ray Films

Assessment of the performance of the X-ray films was made by scanning, mapping, and photometry of the recorded high energy shower events. While ~1000 high energy events were typically observed by visual scanning in one block of an emulsion chamber in the JACEE-12 Antarctic 10-day flight, the number of events recorded by similar selection criteria in the ECT experiment was about 500, mainly because most energetic particles entered from the bottom of the instrument. In fact, the observed number of events above a detection threshold energy of about 3 TeV (sum of gamma ray energies emanating from the interaction) was about 400 events, or 40% of expectation for full-time deep-space experiment. Most primaries entered the bottom of the chamber, having interacted in the material of the bottom of the shuttle bay, in the MPSS structure and the experiment mounting plate. Interactions occurring at a distance from the emulsion chamber produce diffuse showers that are not detected except at very high primary energies.

The scanning and analysis on this point were made at the UAH Cosmic Ray Laboratory by using an in-house designed CCD photometer. A sample of events are shown in a photograph of a flight X-ray film where several high energy cascades are clearly visible. The background darkness discussed in the following section appear as a general gray field in the X-ray films, as shown in a TV picture (Fig. 10a).

All 32 layers of x-ray film in the calorimeter were scanned. The scanned events were projected onto a single map, which gave the direction and incoming zenith angle of each event at a glance, as shown in Fig. 10b. Using this event-map and x-ray films, each event spot on x-ray films was measured for optical density (D_{shower}). The background density (D_{bg}) is also measured around the shower event. The optical density where a shower is located is

$$D = D_{\text{shower}} + D_{\text{bg}}, \text{ where} \quad (1)$$

$$D \equiv \log_{10} I_0/I, \quad (2)$$

and I_0 and I are incident and transmitted light intensity measured by a photometer. These measured data at various radiation lengths in the emulsion calorimeter were plotted as a function of the material thickness (t). Automatic shower fitting for each event was performed at NASA MSFC. Fig. 11 illustrates four such examples. For each shower spot, both D and D_{bg} were measured. To eliminate contamination from the shower in the D_{bg} measurements, the D_{bg} measurements were performed at about 1 cm away from the shower. All the D data for the shower events thus include two values of D : D_{shower} and D_{bg} . Only the D_{shower} 's are used for shower curve analysis. The maximum optical density of an event (D_{max}) is the D_{shower} value at the maximum point in the fitted curve (shower maximum). It is approximately proportional to the number of shower electrons (N_e), and it can be related to the total shower energy as a power function of the shower energy ($\Sigma E\gamma$):

$$D_{\text{max}} \propto [\Sigma E\gamma]^{0.8}. \quad (3)$$

The world-wide convention of equation (1) that defines the shower density D_{shower} from directly measurable film density D and background density D_{bg} is approximate and valid only for low optical density measurements. It is increasingly inaccurate for higher energy events that have high D and/or D_{bg} values. This is because the quasi-linear response function of the Optical Density

of X-ray films gradually deviates from linearity at very high electron densities (ρ), and ultimately saturates to the asymptotic density (D_0),

$$D = D_0 \left(1 - \frac{1}{1 + \alpha\rho}\right), \text{ where } \alpha \text{ is a constant representing silver grain size} \quad (4)$$

High energy events that have high optical density, $D_{\text{shower}} > 2$, in high background-density exposures are subject to corrections corresponding to the exact definition of the subtraction formulae (6) for the optical density of the shower, D_{shower} . The electron density of the shower (ρ_{shower}) and background (ρ_{bg}) have to be used in subtracting the background density from the electron density (ρ_{observed}) observed in X-ray films. The correct electron density and the optical density of the shower at all ranges of the optical density are :

$$\rho_{\text{shower}} = \rho_{\text{observed}} - \rho_{\text{bg}} = \frac{D_0 (D - D_{\text{bg}})}{\alpha (D_0 - D)(D_0 - D_{\text{bg}})}, \quad (5)$$

and

$$D_{\text{shower}} = D_0 \left(1 - \frac{1}{1 + \alpha\rho_{\text{shower}}}\right) = \kappa (D - D_{\text{bg}}), \quad (6)$$

where

$$\kappa \equiv \left[\left(1 - \frac{D_{\text{bg}}}{D_0}\right) - \frac{D_{\text{bg}}}{D_0} \left(1 - \frac{D}{D_0}\right) \right]^{-1}. \quad (7)$$

The ECT experiment is the first to recognize the limitation of the approximate formulae (1) in high density environment. We note here that the exact formulae (6) should be used in any future space experiment where background density D_{bg} is not small. For example, for a film with a background $D_{\text{bg}} \sim 1.0$, the correct value of D_{shower} (from equation 6) may equal 3.5, while the value from equation (1) is $D_{\text{shower}} \sim 3.0$.

4. b):1. High Energy Event Detection and Energy Spectrum.

A total of 383 events was measured by photometric shower densitometry with the selection criterion that the event must have more than 6 layers of the D_{shower} values above the minimum set value, $D_{\text{shower}}(t) > 0.15$. The average number of the events detected with the same criterion for the Antarctic 10-day balloon flight (JACEE-12; 1994) was 864 events. This criterion approximately corresponds to events with the shower energy greater than 3 TeV (primary energy ~ 12 TeV for protons, 40 TeV for irons). The detected events have various zenith angles ranging from 0° to 87° . The spectrum analysis was made only for events with the zenith angle from 0° to 80° , as the D fit for events with zenith angles from 80° to 87° was relatively poor, due to the fact that some of these data were at the edges of X-ray films and there were some uncertainties in D_{bg} data at the edges when measured at the point away from the shower spot. The raw ECT data on the D_{max} differential distribution is shown in Fig. 12a. The integral D_{max} spectrum of high energy cosmic rays from measured events is (Fig. 12b):

$$I(> D_{\text{max}}) = 1005 (D_{\text{max}}/0.1)^{-1.88 \pm 0.096}. \quad (8)$$

The integral energy spectrum $I(>\Sigma E\gamma \text{ (TeV)})$ can be obtained from the D_{max} spectrum by using the relationship, $D_{\text{shower}} \propto [\Sigma E\gamma]^{0.8}$,

$$I(>\Sigma E\gamma \text{ (TeV)}) \propto (>\Sigma E\gamma)^{-1.50 \pm 0.08}. \quad (9)$$

The ECT result formulae (9) is consistent with the all-particle energy spectrum for high energy cosmic rays observed on several JACEE balloon flight experiments, namely:

$$I(>\Sigma E\gamma \text{ (TeV)}) \propto (>\Sigma E\gamma)^{-(1.45 - 1.55)}. \quad (10)$$

This close correlation between measured spectral indices of the gamma-ray inelasticity for the space-flight and balloon flight data confirms the spectrographic capability of the emulsion chamber for high energy cosmic rays. Although a large part of the flight time was earth-facing, and the majority of high charge events interacted with the materials of the shuttle bay floor, the emulsion chamber recorded and identified most of them as interactions originating outside the chamber. Those inversely-developing shower events were degraded in the detectable shower energy ($\Sigma E\gamma$) due to spreading of showers in the path between the vertex (cargo bay floor) and the ECT calorimeter. The inversely-developing events in the integral energy spectrum are reduced in intensity by the reduced ($\Sigma E\gamma$) value which were measured within a finite photometric slit size ($250 \mu\text{m} \times 250 \mu\text{m}$). The loss in the intensity in such a raw (uncorrected) energy spectrum was about 60%, if compared with the prediction for the full-time deep-space flight (~ 1000 events).

Low energy cosmic ray protons ($E < 10 \text{ TeV}$) and shower electrons were major components of the track background in the emulsion chamber. In spite of the fact that a large portion of cosmic rays entered the ECT chamber after passing through the materials of the cargo bay floor, secondary tracks (leading cosmic ray particles, fragments, produced mesons, and showers) were still at relatively high energies, and were not absorbed by these materials. They were accompanied as “inversely developing showers” as shown later in Fig. 19. Hence, the total background intensity due to cosmic rays experienced for the inverted exposure of ECT is approximately equivalent to a 14-day deep-space orientation flight, at least in the majority of the bulk of the chamber.

The recognition of this fact is important as a preamble in the evaluation and extrapolation procedures of the Emulsion Chamber Technology in space for future long-duration orbital flights.

4.b):2 Background Analysis

4.b):2.1 Background Density, Position Dependence and Fluence

The background optical density was in the order of 2.4 in the central part of the x-ray films when a conventional, regular development method (20°C isothermal) was adopted. That of the Antarctic, long-duration balloon flights (10 days) in the similar period (JACEE-12) was 2.2. Both ECT and JACEE-12 received the highest cosmic ray background intensity at the solar minimum period, when the geomagnetic cut-off was the lowest and the lowest energy cosmic ray particles entered the detectors without magnetic rejection. These values of the background densities are approaching the limit of efficient use of X-ray films. The actual films were processed with drastic reduction of background density to $D = 0.2 \sim 0.6$ by a new, low-temperature method, described in the next section.

The background density of x-ray films depends on the position of the film in the emulsion chamber, as a natural consequence of different fluence at different location. The D_{bg} data in films at various depth in the calorimeter are shown in Figs. 13-1 through 13-8, where all the edges indicate higher values of D_{bg} . Because radiation-belt protons would stop within materials less than 20 g/cm^2 , the mid-part of the chamber received high energy cosmic rays and cascade electrons but much less trapped proton radiation; while the edges and the upper portion of the chamber received more proton background and exhibit higher darkness on x-ray films. This can be clearly seen in these figures.

A comment is due for both ECT and JACEE-12 films: the edge density is higher than that of the central area, due to slow protons and soft components that stopped within the chamber. This enhanced darkness in ECT x-ray films at the edges is shown in Fig. 13. These edge darkness values were higher than $D = 2.5$ if processed by a regular development method, causing difficulty using the normal technique with naked eyes and a regular-luminosity light-box. While the use of a high luminosity lamp and a scanning densitometer still allows analysis of these high-density x-ray films, the low-temperature development reduced these darknesses to $D \sim 0.5$, and the analysis was made easy using the standard eye-scanning method.

The absolute value of the D_{bg} by the UAH's CCD photometer was calibrated by the PMT photometer at the NASA/MSFC prior to the STS-62 flight. The uniform background and shower are different in calibrations, because the shower has lateral structure and the CCD and PMT have different saturation functions at high densities. Calibration with uniform density (wedge) is given in Fig. 14a, and that for showers, in Fig. 14b. We use this (Fig. 15a) internal calibration for the general discussion on background endurance in (a)-1 and in other sections. The relationship can be approximated for showers as $D_{CCD} = 1.55 D_{PMT}$ (up to $D_{PMT} < 1.0$). Densities of the high density shower events were measured by both CCD and PMT. On the other hand, the relationship for uniform background is approximated by a 5-th order polynomial (dotted line). The PMT saturates at $D_{PMT} = 4.2$, while CCD saturates at $D_{CCD} = 2.2$ ($D_{PMT} \approx 3.3$). The fluctuations of the photometry for both CCD and PMT were $\sigma = 0.05 \sim 0.06$. Throughout this report, we will omit this error value for simplicity. The D_{bg} values cited in the following descriptions are all those of the CCD measurements, unless otherwise specified.

The thickness (t) dependence of the darkness for the ECT x-ray films is shown by the darkness data in Figs. 13. The Antarctic balloon data, on the other hand, indicated a gradual increase of D_{bg} with increasing material thickness in the calorimeter, as a result of the cascade development of electron showers in the lead calorimeter. (The balloon flight detector did not receive trapped-belt proton radiation.) However, the ECT chamber received all the orbital radiation particles (Fig. 14c) in more complicated manner. When compared with the balloon flight data, the following was observed:

- (1) The (t) dependence of the D_{bg} was quasi-symmetric with the highest values in the center of the calorimeter. The ECT was exposed to cosmic rays and radiation mainly with opposite orientation of the field-of-view to that on balloons.. Consequently, the D_{bg} does not monotonically increase toward the bottom of the chamber. Detailed transport calculations have not been performed to see if this result can be replicated.
- (2) The increase and an eventual decrease of the D_{bg} with increasing depth was more pronounced than the similar fluence data from emulsions and CR-39 that measured the

This observation might offer a plausible explanation to account for the enhanced depth-dependence of x-ray film data, although we cannot quantitatively fully understand the ECT flight data at this point. The analysis is complicated by the mixing of different radiation profiles due to the mixed profiles of the shuttle orientation. The ECT data of D_{bg} in the central location of the x-ray films are shown in Table X1. Compared with this table is the data from the emulsions measured at the center of each plate (Table X2).

Table X1. Three-dimensional data summary of the X-ray film optical density.

Depth (g/cm ²)	C01	C04	C12	C15	C18	C21	C33
A	0.69 ± 0.14	0.59 ± 0.17	0.53 ± 0.16	0.55 ± 0.13	0.50 ± 0.17	0.53 ± 0.16	0.59 ± 0.13
B	0.73 ± 0.13	0.62 ± 0.16	0.59 ± 0.14	0.59 ± 0.13	0.55 ± 0.15	0.55 ± 0.15	0.59 ± 0.13
C	0.63 ± 0.16	0.59 ± 0.17	0.51 ± 0.16	0.51 ± 0.13	0.50 ± 0.16	0.53 ± 0.16	0.58 ± 0.13

A: Center of the left side, B: Center of the film, C: Top edge at the left side.

Table X2. Density of Fog, Grains, and Tracks (at center of the plate).

Vertical Depth	2.29 g/cm ²	33.17 g/cm ²	113.41 g/cm ²	Ground Control
Plate Number	P-03	P-69	C-35	#1
Fog/1000μm ³	1.61 ± 0.14	1.49 ± 0.14	1.71 ± 0.15	1.24 ± 0.13
Grains/100μm	26.09 ± 4.20	30.93 ± 2.96	31.37 ± 2.80	32.42 ± 1.92
Tracks 10 ⁵ /cm ²	4.30 ± 0.16	3.56 ± 0.36	2.53 ± 0.27	0.418 ± 0.19

4. b)2:2 Background Analysis : Isothermal, Low-temperature Development Method Required for Space Flight X-ray Films

When the background density is very high (such as $D_{bg} > 2$), visual contrast in x-ray films becomes very poor for shower detection, and an undesirable saturation of the linear response curve of x-ray films on the shower energy becomes significant. $D_{bg} < 2.0$ is recommended for efficient scanning and preservation of a linear response of D_{max} for energy determination. Considering the possibility of much higher background density, as would result by exposing the EC on Space Station for 1/2 to 1 year, we experimented and established a new, low-temperature development recipe for the ECT experiment (5 °C isothermal for 3 minutes). Table X3 shows a comparison of conventional recipe and the ECT's new recipe.

Table X3. Comparison of recipes for X-ray film development

Low-Temperature Method	Development	Stop	Fix
Chemicals	Konidol	Acetic Acid 3%	Konifix
Temperature °C	5	5	5
Duration (minutes) -variable	1 ~ 7	1 ~ 3	30 ~ 40

Conventional Method	Development	Stop	Fix
Chemicals	Konidol	Acetic Acid 3%	Konifix
Temperature °C	20	20	20
Duration (minutes) -variable	5 ~ 20	1	20

The new ECT prescription seems to respond well to the particle-background density problem posed by long-duration space flights, by reducing the D_{bg} from 2.1 to 0.5 without compromising the number of detectable events. (We used longer development time of 5 minutes in the actual development of the ECT X-ray films, with the average D_{bg} value of 0.2 by CCD and 1.1 by PMT.) The method works by reducing *the size* of the developed silver-halide crystal without much reducing *the number* of latent image-grains. Since D_{bg} varies with the background track density, N_e (bg) approximately according to:

$$D_{bg} \propto \log N_e \text{ (bg)}, \quad (11)$$

we project that the effectiveness of x-ray films in recording and analyzing events will be maintained up to backgrounds of ~80 times ($10^{(2.4-0.5)}$) that of the ECT exposure. We conclude that, with suitable adjustment of development procedures, chambers can be effectively deployed (in a similar orbit) for up to $(80 \times 15) = 1200$ days.

At the higher orbit (400 km) and inclination (57°) expected for the International Space Station, the background density can be about 3 times higher than the STS-62 orbit (300 km, 39°), and the maximum useful duration would be about 400 days. This number is subject to the nature of the background. The number quoted here is for trapped belt radiation, which affects the validity of x-ray films in the edges and at the shallow depth in the emulsion chambers.

4. b):2.3 Background Analysis: Materials Compatibility Studies

Aluminum and other "active" metals have long been known (ref 5) to have potentially damaging effects on silver bromide emulsions, and emulsion lore is rife with stories of plates and pellicles ruined by contact with or proximity to such metals. The JACEE Collaboration in 10 balloon flights had avoided that problem by constructing the emulsion containers entirely from non-metallic materials such as rubber sheet, polymethylmethacrylate (lucite) and plywood. The rigors of rocket launch, and the requirements of demonstrably safe mechanical confinement and attachment, resulted in the basic container design being a hermetically sealed aluminum box. While this conferred some advantages, such as more accurate plate positioning and maintenance of constant humidity in the plate materials, it did require a focused effort to assure compatibility of materials with emulsion gel plates and x-ray films. These detectors must be able to be stored in the flight housing for 1 year without significant degradation.

A program was devised and carried out in which small pieces of the detector materials were exposed to the box construction materials for various periods from 1 to 18 months. Of

principal concern was Al. Tests were conducted with bare Al and with various kinds of surface coating on the metal. Tests were conducted both with the dissimilar materials both in direct contact, and in close proximity within small sealed chambers.

Table X4 lists the materials and conditions of exposure of films. Diagnosis is defined by the words "Normal" or "Damaged".

Table X4. Various tested materials, Optical Density (PMT) and physical conditions.

<i>Hermetic</i>	Sample 1 (1.0 months)	Sample 2 (2.5 months)	Sample 3* (4.5 months)	Sample 4 (18 months)
Controls	1.46	1.50	0.91	2.02
with Krylon	1.47	1.46	0.89	-
with Epoxy	1.45	1.58	0.94	-
with Aluminum	1.38	1.56	0.94	-

All samples were normal conditions; * short development processing; - Data not available

Table X5 shows the optical densities of the test films. The sample names, 1, 2, 3, 4, and F, correspond to the duration of the exposure for the test, 1.0 month, 2.5 months, 4.5 months, 8.5 months and 18 months, respectively. The sample 4C was for 8.5 months.

Table X5. Optical Density (PMT data) of the ground control X-ray films.

<i>Material/Period</i>	<i>Sample 1</i>	<i>Sample 2</i>	<i>Sample 3</i>	<i>Sample 4</i>	<i>Sample 4C</i>	<i>Sample F</i>
Control	1.57	1.55	1.55	2.02	-	1.38
Aluminum#	2.52@	2.19@	2.20@	3.48@	2.07@	1.78@
Anodized Al.#	1.76#	1.80#	1.93#	2.17#	1.69#	1.52#
Alodined Al.#	1.69##	1.55##	1.57##	1.97##	1.98##	1.65##
Black Lucite	1.45	1.44	1.49	1.55	1.92	1.80
Clear Lucite 1/4"	1.43	1.51	1.52	1.80	2.02	1.47
Si Rubber	1.49	1.54	1.49	1.64	1.78	1.52
Clear Lucite 1/16"	1.67	1.62	1.61	1.72	1.87	1.55
Viton O-ring	1.61	1.60	1.56	1.78	1.68	1.45
Gold Plated AL.	1.41	1.50	1.50	1.72	-	1.50
Parafilm	-	-	-	1.73	-	1.66
Anodized Al. painted 1	-	-	-	1.84	-	1.50
Anodized Al. painted 2	-	-	-	1.78	-	1.49
Pb (cleaned)	-	-	-	-	-	1.78
Pb (painted)	-	-	-	-	-	1.65

@: All contact pieces were damaged, progressively worsened; non-contact part was fogged but not damaged.

#: All contact samples were somewhat damaged; non-contact sample were not damaged.

##: Some contact samples were damaged; non-contact samples were not damaged.

All other samples without @, #, and ## symbols were in "Normal" condition.

Aluminum and alodined aluminum were shown to destroy emulsions and X-ray films on contact, while anodized aluminum was inert. Anodized aluminum was selected for the space flight emulsion chamber based upon these material compatibility tests. The actual shuttle flight result confirmed the ground test results. It is reasonable to extrapolate further, based on the ground tests and the ECT flight, that all the materials in the flight emulsion chamber in a ECT pressurized vessel should be safe for long duration space flights at least up to 18 months.

4.c) Emulsions

Measurements of the high energy cosmic ray events, background track density, chemical fog, and grain density were performed using high magnification microscopes. Similar measurements were performed with ground control unit and balloon-borne emulsions to compare the quality and capabilities with the space-flight emulsions.

4.c):1 Quality of Tracks (Examination by Berriman-Curve Test)

The contrast for track recognition in emulsions was excellent and clear track measurements were achieved. Grain density relative to the fog density is a measure for evaluating the quality of the track recognition contrast (Berriman curve Fig. 15). "Excellent" quality of emulsions are indicated by a domain above the solid curve in the figure, while the dotted curve represents the "good recognition". They were measured at the central part of the emulsions, because the majority of the emulsion measurements for cosmic rays depend on the quality in the central part where the event tracking will be made. All the ECT emulsions turned out to be "excellent contrast" as demonstrated in Fig. 15 (Berriman curve) and Fig. 16 (photograph).

Table E1 provides the measured densities of grains and fogs, as well as the background track densities. In average, the grain density of the ECT flight emulsions for relativistic, minimum-ionizing, $Z = 1$ particles was 29.5 ± 1.8 grains/100 μm , while the fog density was 1.60 ± 0.08 fogs/1000 μm^3 , while those of the ground-control emulsions for the same period were 32.42 ± 1.92 grains/100 μm and 1.24 ± 0.13 fogs/1000 μm^3 , respectively. The grain density of both materials are similar within the statistical errors, while the fog density is clearly enhanced by 29% (to 3σ level) in the flight emulsions. This difference is small enough to assure the high quality of flight emulsions.

Table E1. ECT background measurements (on emulsion properties).

Plate Number	Location from the top (g/cm ²)	Fog density (fogs/1000 μm^3)	Grain density (grains/100 μm)	Background (10 ⁵ tracks/cm ²)
P-03*	2.29	1.61 ± 0.14	26.09 ± 4.20	4.30 ± 0.16
P-69**	33.2	1.49 ± 0.14	30.93 ± 2.96	3.56 ± 0.36
C-35**	113.4	1.71 ± 0.15	31.37 ± 2.80	2.53 ± 0.27
Control**	GROUND	1.24 ± 0.13	32.42 ± 1.92	0.42 ± 0.19

* measured at 1 cm from the edge of the plate.

** measured at the center of each plate.

The ECT in-flight temperature set-point was selected at 20°C. This is near the upper limit of safe working temperatures for emulsions (30°C maximum). Emulsion fog is often induced by high temperature and by some active chemicals in the environment surrounding the emulsions. It is possible that the small observed difference in fog densities is mainly due to thermo-chemical fog

induced during some part of the transportation and storage period when several days of high temperature (21 - 22.5°C) were recorded for the flight emulsions. The ground-control unit did not experience such high temperatures. **Table X5, Table E1 and Fig. 15** show a time-and-temperature dependence of fogs in emulsions. In the following discussion we should emphasize that the fogging observed in the ECT plates was a small but observable effect that did not affect data retrieval to any significant degree.

The compatibility test (described below) indicated a very slow increase of fog as a function of the increasing storage period at room temperature ($\approx 17^{\circ}\text{C}$). In the anodized aluminum box in which the ECT emulsions were flown, the fog increased from only 5% to 10% for a storage duration of 1 month to 18 months over the hermetically sealed storage emulsions (contact with Lucite plates). The ECT result implies that the observed increase of fog over the ground-storage materials was not likely to have originated from the chemical reaction with the materials in the ECT box in the space environment, but it is more likely due to the increased temperature during the periods (Jan 6 -13; Jan 26 - 30; Feb. 1 - 5, 1994) prior to the STS flight (**Fig. 17**).

Although the analysis of fog indicated with reasonable likelihood that high temperature during ground storage/transportation was responsible for an increased fog of about 30%, a further question remains: whether there was any combinatory fog increase due to other materials in the chamber, interacting at higher temperatures than that of our compatibility tests. This examination will be necessary to fully guarantee a limited fog-increase for a very long-duration space flight, and it must be addressed here as a further requirement of a ground test.

4.c):2 Tracing of High Energy Tracks/Showers in Emulsions

The flight emulsions provided high visibility for all individual tracks including minimum ionizing tracks. Very clear shower tracks were photographed from emulsions showing excellent quality of event recording as demonstrated in **Figs. 18 and 19**. A "normal shower development" is shown in **Fig. 18**, which is an event entered into the chamber from space in deep-space flight orientation period. An "inverse shower development" is easily identified by the unique signal of the inverse development of the lateral spread. Shown in **Fig. 19** is an example where the event produced a cascade shower from the bottom to the top of the chamber after entering and interacting with the cargo bay during the Earth-observing orientation of the STS-62.

A primary iron nucleus track and two interaction vertices in the emulsion chamber were photographed in **Figs. 20 - 22** with the highest magnification of x 100 objective lens. General (x 20) image of emulsions were compared in **Fig. 23** between an ECT emulsion and a long-duration balloon-borne emulsion, where similar excellence of track qualities are observable.

Showers were traced from the lower part of the calorimeter upward into the target and primary modules. This event tracing is expected to be subject to interference by copious background tracks at high background density. Despite the density in the order of $10^6/\text{cm}^2$ in the ECT emulsions, there were no significant difficulties in tracing events upwards even when the "jet" structure of the event becomes thin and small in the target module.

The ECT and long-duration balloon-borne emulsions (JACEE-12, 1994; JACEE-13, 1995) have the highest background level among all the past space-flight emulsion chamber experiments. It is not straightforward to assess what density of the background will prevent efficient tracing, because we do not have much experience with emulsions that have much higher background density than the ECT or JACEE-12. Nevertheless, some assessment is possible.

Emulsion is a three-dimensional micro-photographic device. The vertical focus is as sharp as $1\text{ }\mu\text{m}$, and the lateral resolution of tracks is as good as $0.1\text{ }\mu\text{m}$. One minimum-ionizing, relativistic track will form about 30 grains in $100\text{ }\mu\text{m}$ of track length. The visibility can be approximately defined by the average grain distance (r_{bg}) between different tracks versus average grain distance in a track (r_g):

$$r_{bg}/r_g > \lambda, \quad (12)$$

where ($\lambda \geq 1$). Sufficient track recognition with a condition of ($\lambda \sim 2$). The minimum condition to identify a track corresponds to ($\lambda \sim 1$).

A cosmic ray track has an average zenith angle of 45° , and therefore, the average lateral grain separation is $2.36\text{ }\mu\text{m}$, while that of vertical is also $2.36\text{ }\mu\text{m}$. This chain of grains (blobs) belonging to a single track can be recognized as a track when other grains from other tracks overwhelm and confuse the recognition of a single particular track. When one has $10^6/\text{cm}^2$ track density in emulsions (ECT) the average track separation (r_{bg}) is in the order of 14 microns, satisfying the above ratio $r_{bg}/r_g > 1$ by a factor of about 5. If the background cosmic ray tracks (not those of trapped-belt radiation protons which will be absorbed in emulsion chamber) accumulates up to 35 times more than the ECT track density, the single ionizing track recognition becomes very hard as the r_{bg} becomes as large as r_g . This is true for the high-sensitivity emulsions (Fuji ET-7B). Hence, our conclusion should be that the single ionizing track can be traced with increasing difficulty in high background track densities up to 3.5×10^7 particles/ cm^2 , and the ECT's 14 day flights had only 1/35 of this limit. The limit may be 35×14 days = 490 days on orbit, provided that the emulsion chamber is large enough to absorb most of slow proton background coming from the radiation belts. The edges and the very top portion of the emulsion chamber record slow protons before their stopping, and the visibility of a single track therein will be poorer in these edge regions.

Concerning traceability in low sensitivity emulsions (Fuji ET-6B) for nuclei with charge of helium or larger; the same argument applies with different parameters. The r_g of a Helium track in ET-6B is $2.53\text{ }\mu\text{m}$. Because ET-6B records grains of background protons as few as 7 grains/ $100\text{ }\mu\text{m}$, they can be completely ignored. The background tracks to be considered are those of helium and $Z/\beta \geq 2$. They are less than 30% of cosmic rays. Therefore, from the consideration of r_{bg} , the limit of track recognition will not be reached until exposures of 1,420 days. This limit with the low-sensitive emulsions (ET-6B) will remain close to this value even when radiation-belt protons increases in a very high orbit (500 - 1000 km), so long as the main detector part where the self-absorption of the emulsion chamber effectively works is concerned.

For heavier tracks ($Z > 2$), the average grain separation in low-sensitivity ET-6B emulsions is $1.12\text{ }\mu\text{m}$ for lithium ($Z = 3$), and $0.28\text{ }\mu\text{m}$ for Carbon ($Z = 6$). Acceptable track densities for them are very high, and the limit of exposure duration in space would exceed several years, so long as low-sensitivity emulsions are used.

4.c):3 Track Density in Emulsions

The track density was measured by both manual and automatic microscopes. Track by track identification was easily made by manual visual scanning as listed in Table E1.

The automatic microscope (CUE-2) does not identify blobs and separate grains as belonging to the same track, and gives an order of magnitude larger number of objects in a field of

view. (An advanced algorithm to connect the spatially-separated parts of a track [objects] into an identified single track is being developed, but it has not yet been used in the ECT analysis.)

The Table E2 shows the material-thickness dependence of the object density in the ECT and JACEE-12 emulsion chambers. By the distinct identification of real tracks in the manual scanning, the number of objects in the CUE-2 auto-analysis can be calibrated as 10 times of the number of individual cosmic tracks.

Table E2. "Object" density dependence on vertical material thickness.

Object Density ($10^6/\text{cm}^2$)	P-03 at 2.29 g/cm^2	P-45 at 22.2 g/cm^2	P-69 at 33.2 g/cm^2	C-10 at 50.9 g/cm^2	C-14 at 60.9 g/cm^2	C-18 at 70.9 g/cm^2	C-22 at 80.9 g/cm^2	C-26 at 90.9 g/cm^2	C-30 at 101. g/cm^2	C-35 at 113. g/cm^2
ECT	3.59	4.46	4.70	4.02	4.08	3.53	3.71	1.55	0.99	0.34
JACEE-12	1.84	2.99	4.57	4.53	-	-	3.91	-	-	-

- (data not available at this time at the same vertical thickness)

4.c):4 Linear Energy Transfer (LET) Data

Track grains (blobs) were counted for individual particle tracks with a microscope having x 1,500 magnification. A blob is a clump of grains that are not separable with the 1,500 magnification. More grains are viewed as blobs for tracks that have higher Linear Energy Transfer (LET), and the present results on LET spectrum are lower bounds at higher LET's. Figs. 24 (a) - (c) show LET spectra at the vertical thickness of 2.29 g/cm^2 , 33.17 g/cm^2 , and 113.4 g/cm^2 , respectively.

The intensity of the minimum ionizing tracks does not change much with the material thickness. However, tracks that have $dE/dx > 2 \times$ minimum ionization ($\sim 4 \text{ MeV}/\text{g}/\text{cm}^2$) decrease substantially with increasing material thickness. While the details of the thickness dependence must wait for a full Monte Carlo simulation for quantitative analysis, it can be concluded that the observed profile should reflect absorption of slow protons of trapped belt radiation. This conclusion is also supported by comparison of the LET spectra.

At shallower depths of the ECT and JACEE-12 materials, the LET spectra are dissimilar at high LET's: Relative intensity of high LET particles at shallow material thickness are much more abundant in the ECT experiment than that of the JACEE-12, while those at deeper depths are similar, indicating that the high LET tracks of ECT materials are absorbed in the ECT materials.

For comparison, JACEE-12 data are shown at equivalent thickness (1.5 g/cm^2 , and 70 g/cm^2) in Figs. 24 (d) ~ (e), respectively.

4.d) CR-39 Analysis

CR-39 etchable plastics were separately described in this final report by the University of San Francisco co-investigators. Several independent measurements were performed at the University of Alabama in Huntsville, described below.

Owing to operational problems and increased environmental restrictions on the chemical etching facilities, the etching of the ECT CR-39 has just been completed. We have on hand now in

our lab two sets of CR-39 plates fabricated from material of the same specification (American Acrylics USF-3). One set flew in orbit for 14 days in ECT; the other for 10 days on an Antarctic balloon. A few notable differences were discovered in these two different flight experiments. An assessment of the difference of sensitivity was made by taking a temperature effect into account. Assessment of the difference in physical appearance was made in terms of scattering centers formed by small etch-pits and contaminants, nonetheless, thorough assessment requires additional material compatibility tests of CR-39 at different temperatures and at different atmospheric pressures.

4.d):1 CR-39 Etching, Uniformity, and Quality

Large size CR-39 plates (40 cm x 50 cm) were included in the ECT and regular balloon flight emulsion chambers. They are intended to be used for charge measurements of heavy nucleus tracks, and are also considered for the state-of-the-art coordinate measurements for future analysis of track momentum by using the multiple Coulomb scattering method.

The large-size CR-39 plates exposed in previous experiments were usually cut into smaller segments and etched (20 cm x 25 cm or smaller size) so that they fit the stage of measurement microscopes. The University of Alabama in Huntsville has developed a large stage microscope (50 cm x 50 cm) to analyze a large-size CR-39 plate for coordinate measurements. Correspondingly, the ECT CR-39's were etched in their original shape (40 cm x 50 cm) in a large etching bath at the Naval Research Laboratory. Standard etching recipe was used: 70°C for 24 hours with 6.25N NaOH solution. Two large racks made from stainless-steel wire-mesh were used in a etching bath. The temperature gradient over the entire racks were monitored at various rack positions and was controlled to within $\pm 0.05^\circ\text{C}$.

4.d):2 CR-39 Objects (Etch-pit hole) Data

CR-39 plates were scanned with an automated "object" analysis microscope (GALAI CUE-2 Auto-morphometer). Each plate was measured at the center (Part B) and at opposing edges (Parts A and C). Many parameters were measured automatically during the scanning operations. There were many small etch-pit holes and the measured data of the "area size", which include those of background stopping α -particles, and contamination due to chemical instability of CR-39 surfaces. The ECT plates had a larger population of these small etch-pits relative to those of the JACEE-12 balloon flight materials, although these two experiments used the same CR-39 formulation from the same lot.

The measured "track density" strongly depends on how many of these small etch-pits are contained. Also, "average area-size" similarly depends on them, but in the opposite way: the "average size" becomes smaller when more small-sized contaminants are included. To minimize the effect of small-size contaminants, the auto-program set the minimum sampling value ($60\ \mu\text{m}^2$) for the "object" area-size.

Three different sets of CR-39's were included in the ECT emulsion chambers. The group (I) are 10 sheets of CR-39 (CR1 ~ CR10), used in the emulsion chamber for regular charge measurements of high energy nuclei. The group (II) consists of 6 sheets (B2 - B7) inserted in the calorimeter section, which contained dozens of CR-39 doublets (E04 + E01, E12 + E05, ...) in cut-out slots. The dosimetry and LET spectra in the calorimeter are reported by using these materials in the separate article by the University of San Francisco group.

We measured in several places on the large CR1 - CR5 plates from group (I), and two calorimeter plates of the group (II) that held doublets of (E16-EC) and (E30-E38). The “object density” and the “average area” of objects are shown in **Figs. 25a and 25b**, respectively, as a function of increasing vertical thickness and the location (A, B and C). The central part (B) had less variation in the “object density” than those at edges (A and C). High object density at edges (A and C) indicated gradual decrease with increasing thickness. The average area size for all parts (A, B and C) did not show noticeable differences throughout all the depth for CR1 to CR6. Nonetheless, as remarked previously, these data are still subject to change due to unclear origin of contaminants of small objects. The analysis by CUE-2 auto-morphometry remains uncertain in this regard.

4.d):3 CR-39 Data Comparison with Balloon-borne Environments

Similar analysis was performed for CR-39 plates flown by a balloon (JACEE-12). **Figs. 26a and 26b** show the “object density” and the “average area” from the top (RRP2) to the bottom (RRC22) of the chamber. Only small variations on these data were found as a function of the increasing thickness of the chamber materials. However, the data on the top indicate higher density and average size, which quickly decreased within about 10 g/cm², suggesting absorption of slow nuclei (nearly zero cut-off energy) received near the magnetic south pole in the JACEE-12 circumpolar flight.

The most noticeable difference of etch-pit hole data between the ECT and JACEE-12 flights is the average area size. The ECT data indicates the median of ~ 250 μm², while the JACEE-12 balloon flight data shows that of ~ 400 μm². The observed difference is somewhat puzzling, because they were produced in the essentially same lot and etched at almost the same time by the same NaOH prescription. The cause of this difference is not clear. Nevertheless, we have also recognized another difference between the two samples; namely, the ECT plates were rather milky (having a lot of scattering elements ~ small etch-pits) while JACEE-12 plates were far more transparent (having less scattering centers). The number density of small etch-pits in CUE-2 auto-morphometry in fact supported this observation of the physical appearance.

The observed difference of sensitivity cannot be straightforwardly accounted for by the small difference of the charge and energy spectra of heavy nuclei in orbital flight and circumpolar balloon flight. A large difference (~ 40%) of the average area of large etch-pits is suspected as a possible environmental effect on CR-39 sensitivity during the flight.

It is well known that the sensitivity of CR-39 is strongly dependent on temperature, particularly when they are compared at low temperature (T ≤ 0°C) and at ambient room temperature (~ 20°C) (**Fig. 27**). The former are about 50% more sensitive than the latter case, and the observed difference (~ 40%) is reasonable.

The second difference, the number of small etch-pits or contaminants, can hardly be attributed to the temperature effect only, because the ground control did not show the consistent results to support the temperature effect on this characteristic.

More importantly, these two flights had another physical difference during the flights when the cosmic tracks were recorded. The CR-39 sensitivity has been known to be low in vacuum condition. The latent image requires oxygen for track record to be fixed in polymerized molecular reactions. Because of this reason, the ECT used a pressure vessel (~ 1 atm. of air) to maintain the sensitivity of CR-39. JACEE balloon flights were always carried out with emulsion chambers

enclosed in an air-tight rubber-bag with a one-way passive valve to release interior gas out to environmental low pressure. The JACEE chamber was literally in vacuum when tracks were recorded, and did not benefit from the positive effect of oxygen for fixing the latent image of etch-pit hole. This physical difference on oxygen accompaniment with CR-39 plates can hardly account for either of the differences observed between orbital and balloon flight.

A suspicion exists for possible combinatory chemical-thermal effect on CR-39 plates. We have conducted materials compatibility tests for emulsions and X-ray films. However, no such tests were performed in the ECT experiment. Some materials in the emulsion chambers may be reactive with CR-39 surfaces and can cause irregular contaminants or quasi-etch pit holes of very small kind. Such effect, if it exists, must be activated either by moderately high temperature ($\sim 20^{\circ}\text{C}$) and/or oxygen content. Though it remains speculative, the ECT experiment alone cannot exclude such a possibility. Further material compatibility tests of CR-39 plates at high and low temperatures with and without oxygen must be performed in combination with all the emulsion chamber materials.

5. Conclusions: ECT Flight Data and Projected Feasibility for Long-Duration Space Flights

The ECT experiment provided sufficient materials and data to demonstrate excellent performance of an emulsion chamber for an orbital flight, and to verify the basic design approach for containment and environmental control throughout all flight operations.

The pressure chamber and active thermal control system operated normally throughout shuttle launch, orbital flight, and landing. The pressure was kept at 1 atm all the time and the ECT materials were maintained at $20 \pm 0.1^{\circ}\text{C}$ and 1.0 atm pressure during track registration.

The x-ray films in the emulsion calorimeter registered as many high energy cosmic ray events as expected by the pre-flight calculation using the known cosmic ray flux. The emulsion quality was found to be as good as any balloon flight experiment or ground control unit for recording cosmic ray tracks from protons to iron nuclei, including secondary mesons and cascade electrons.

The outer few cm of the dense emulsion chamber material was found to absorb efficiently the slow protons and electrons from trapped-belt radiation. The track acceptance capacity in emulsions and X-ray films and usefulness for track analyses are found to be limited more by cosmic ray intensity than the radiation dose from trapped belt particles.

The track densities of minimum ionizing tracks allowed an assessment for extrapolated long duration space flight. The ECT emulsion chamber allows track analysis of cosmic ray protons up to 1.34 years of orbital flight. Longer space flights using low-sensitivity emulsions may be useful for up to 3.9 years for helium nuclei, and even longer for carbon and larger nuclei.

The cascade recording by X-ray films was found to be useful for long-duration space exposures, with some adaptations of current method. In particular, a low-temperature under-development method was invented during the ECT experiment, allowing high-track-density exposures of the order of ~ 80 times the ECT flight for useful shower analysis, although the threshold energy will increase accordingly ($\sqrt{80}$), from $\Sigma E_{\gamma} \sim 2$ TeV for 14 day flight to $> \sim 20$ TeV for up to ~ 3.3 years of flight.

The CR-39 solid state track detector was useful in orbital flight, giving excellent LET spectra at various material overburdens. Some differences of the orbital flight materials in ECT chamber from those of Antarctic balloon flight were observed. The main difference, that of sensitivity, was accounted for as due to temperature effect. We should recommend a re-examination of the set-point for the thermal control system for a future flight. A lower set-point in the range of -5°C to 10°C would use less power, and provide a better operational environment for emulsions and particularly CR-39. Additional material compatibility tests of CR-39 are recommended for preparing CR-39 exposure in emulsion chamber for long-duration space flights.

6. References

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EMULSION CHAMBERS

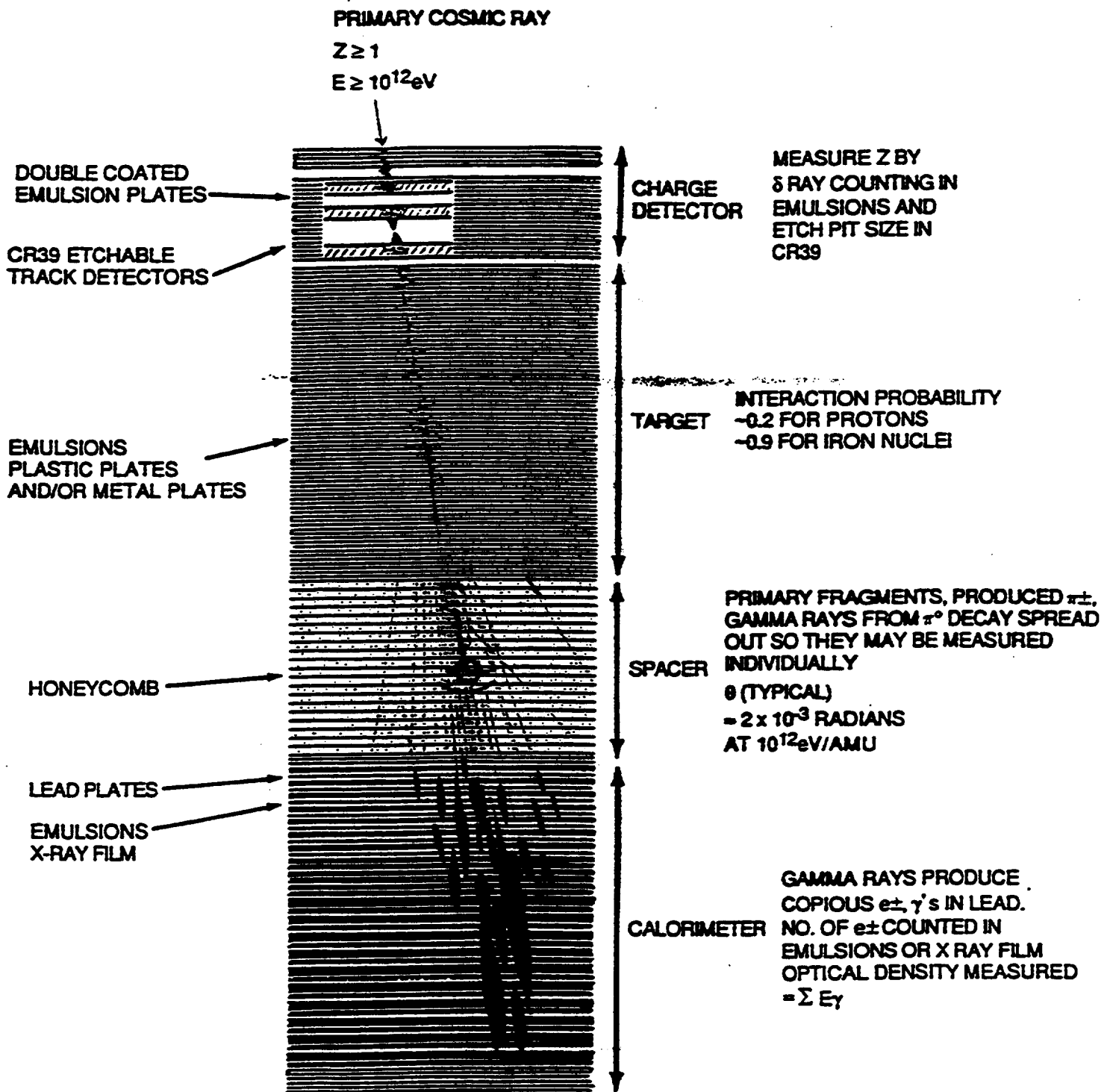


Figure 1

FOIL HEATER ON TOP OF LID
SILVERIZED TEFLON ON TOP OF HEATER

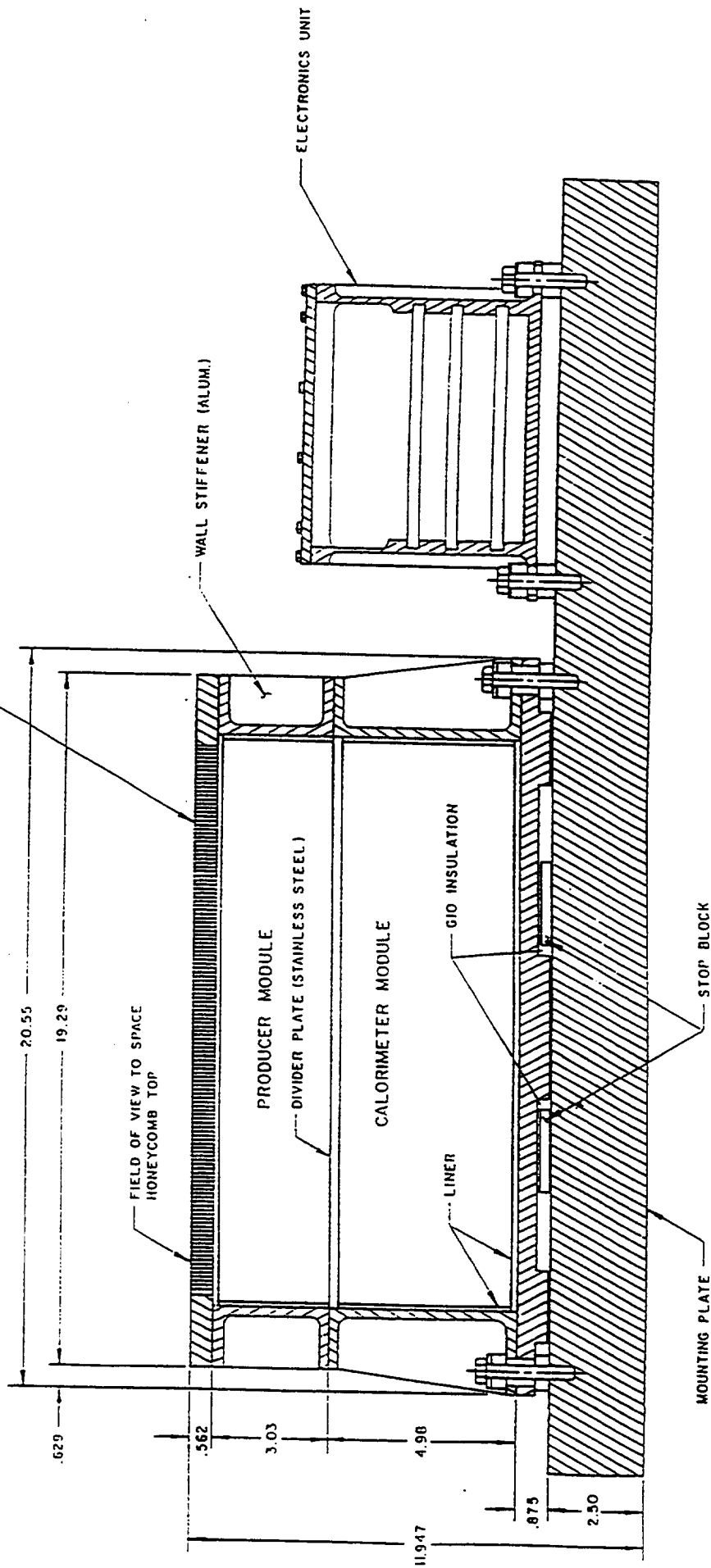


Figure 2

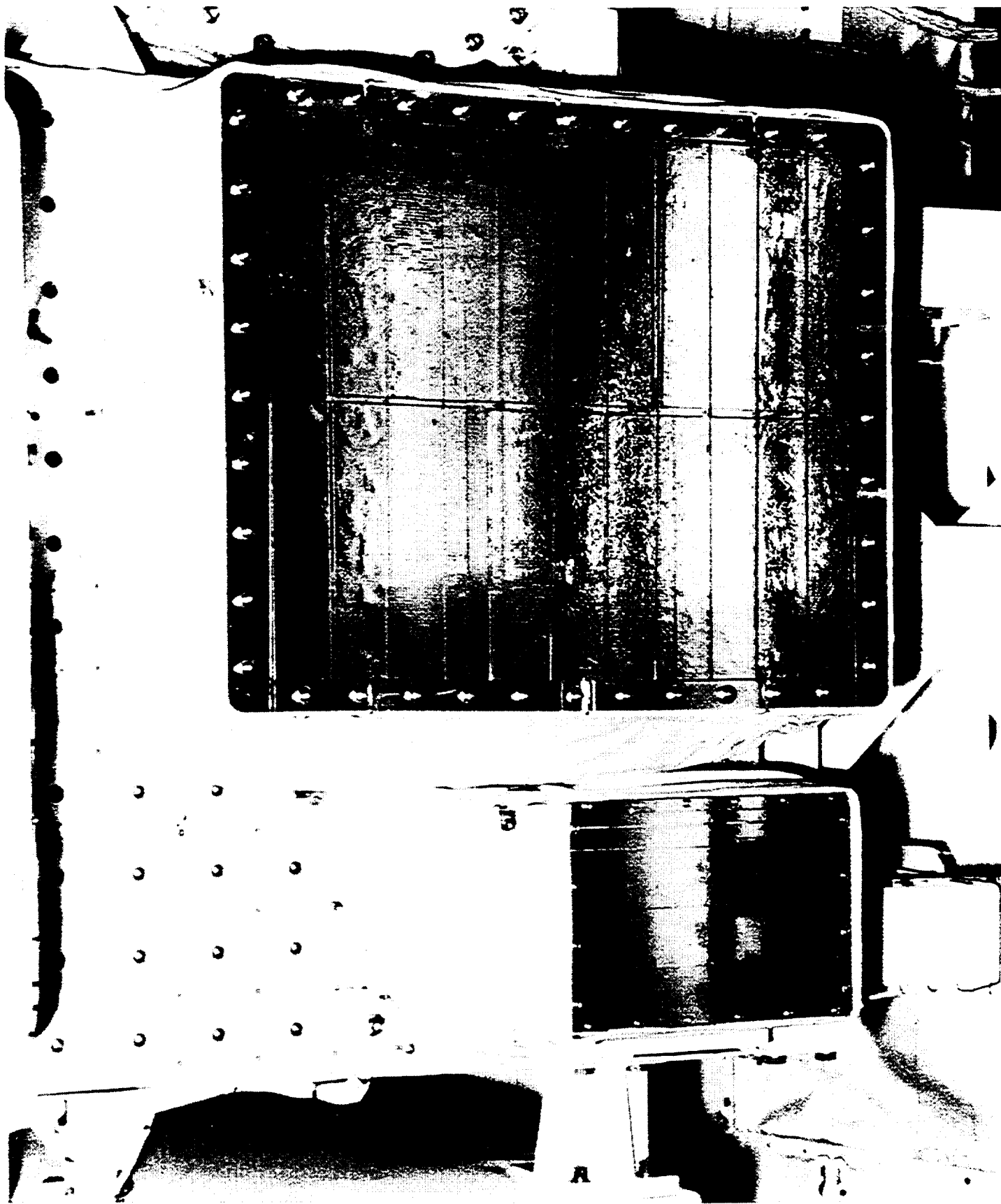


Figure 3

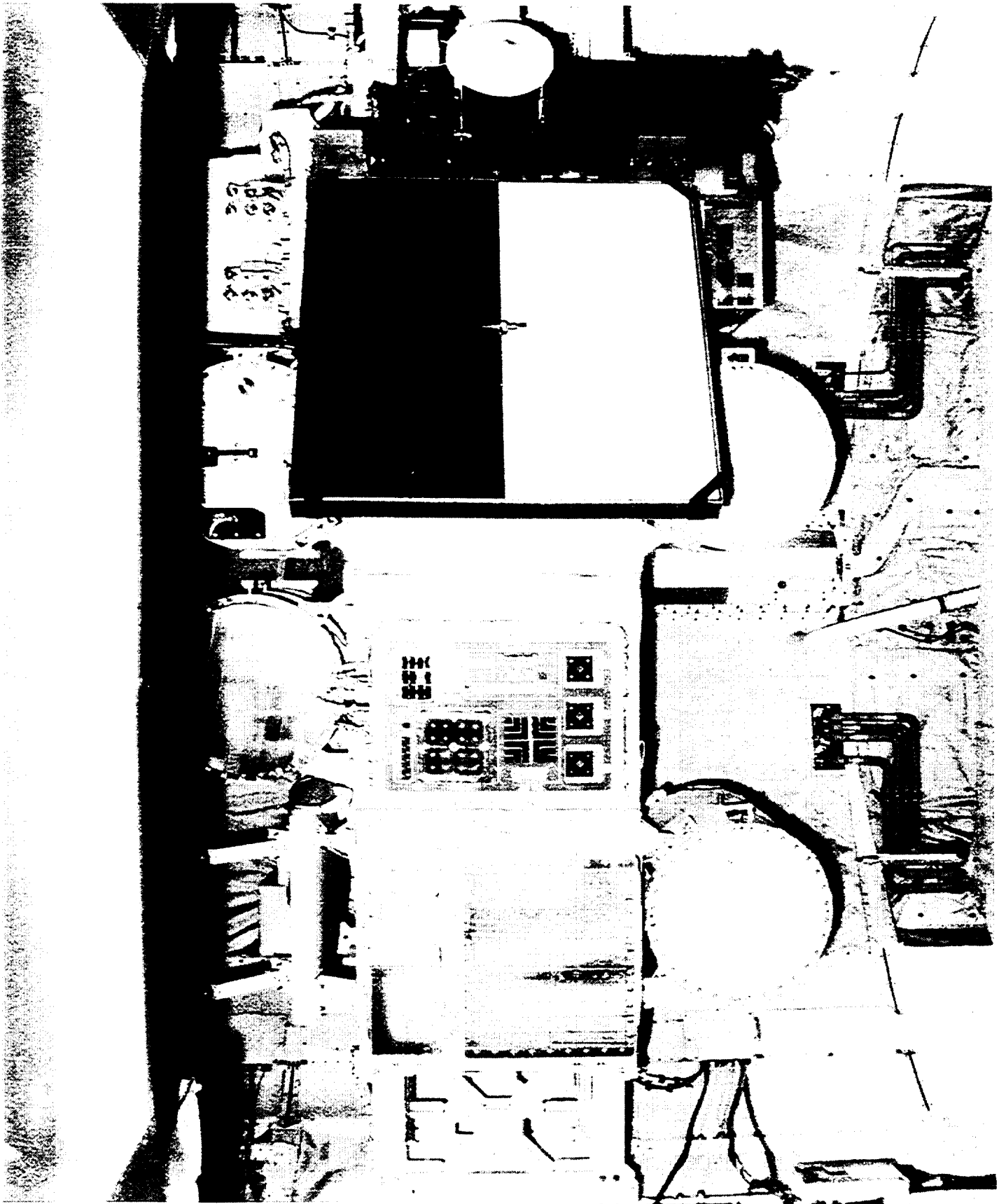


Figure 4

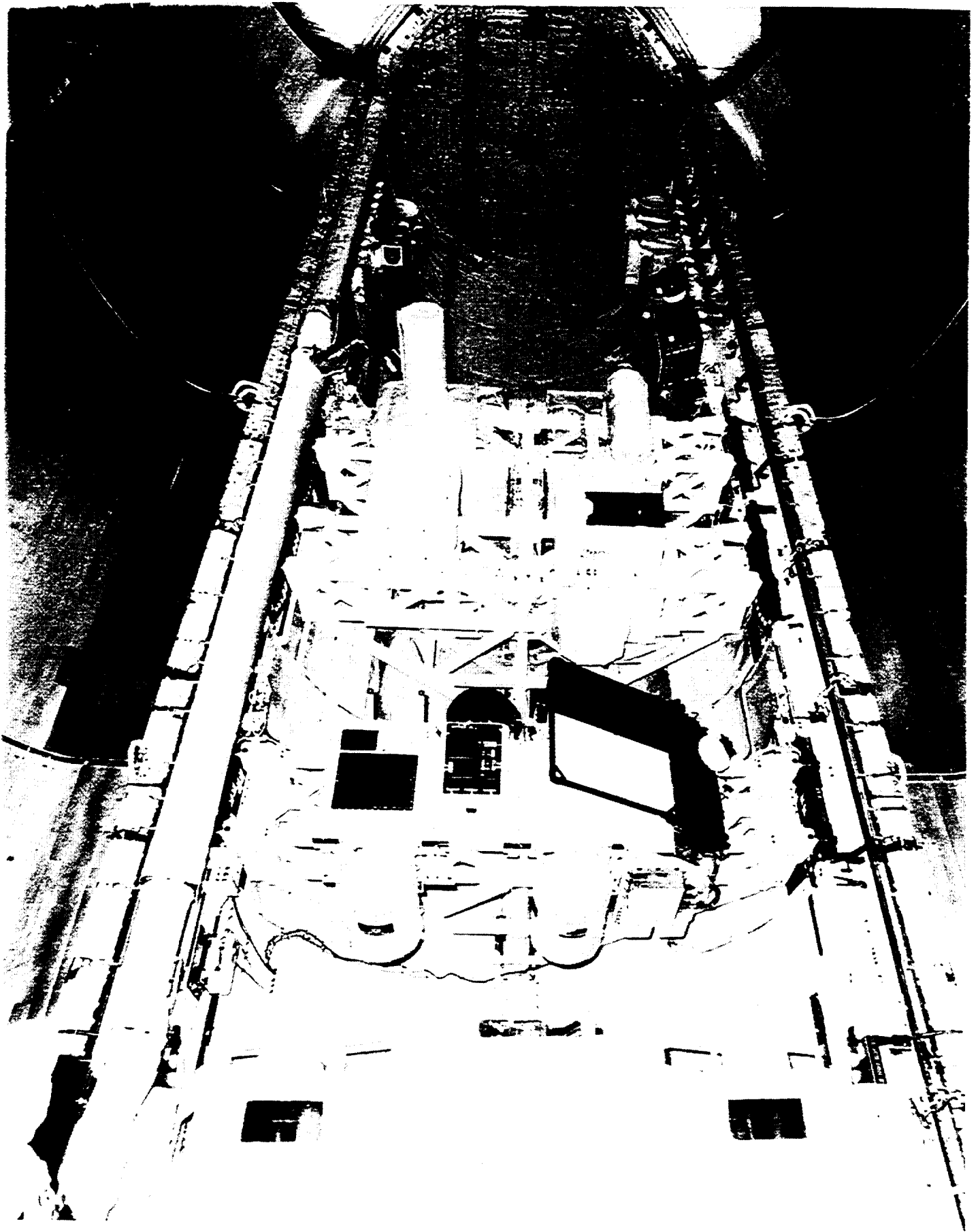


Figure 5

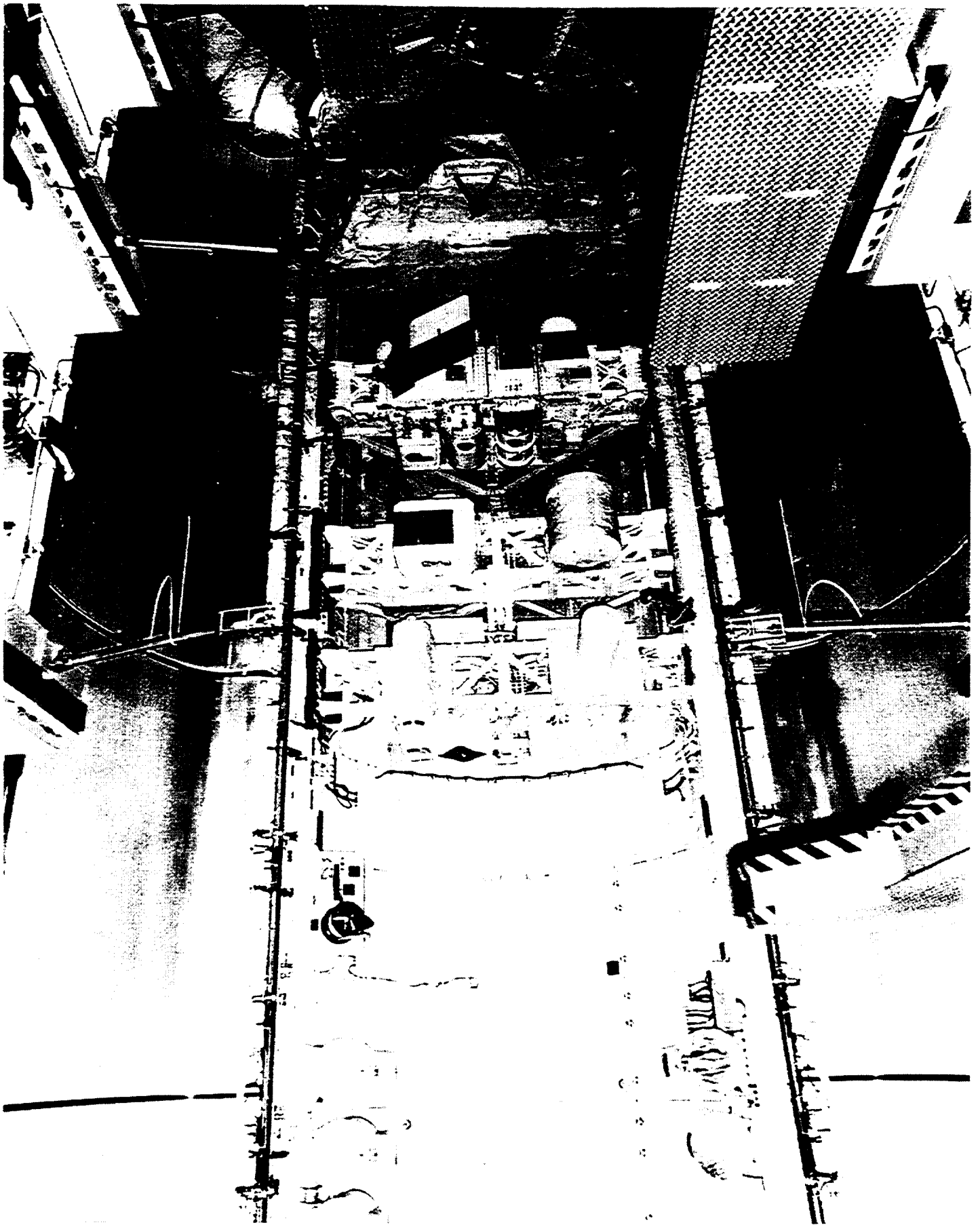


Figure 6

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Forward MPES Configuration

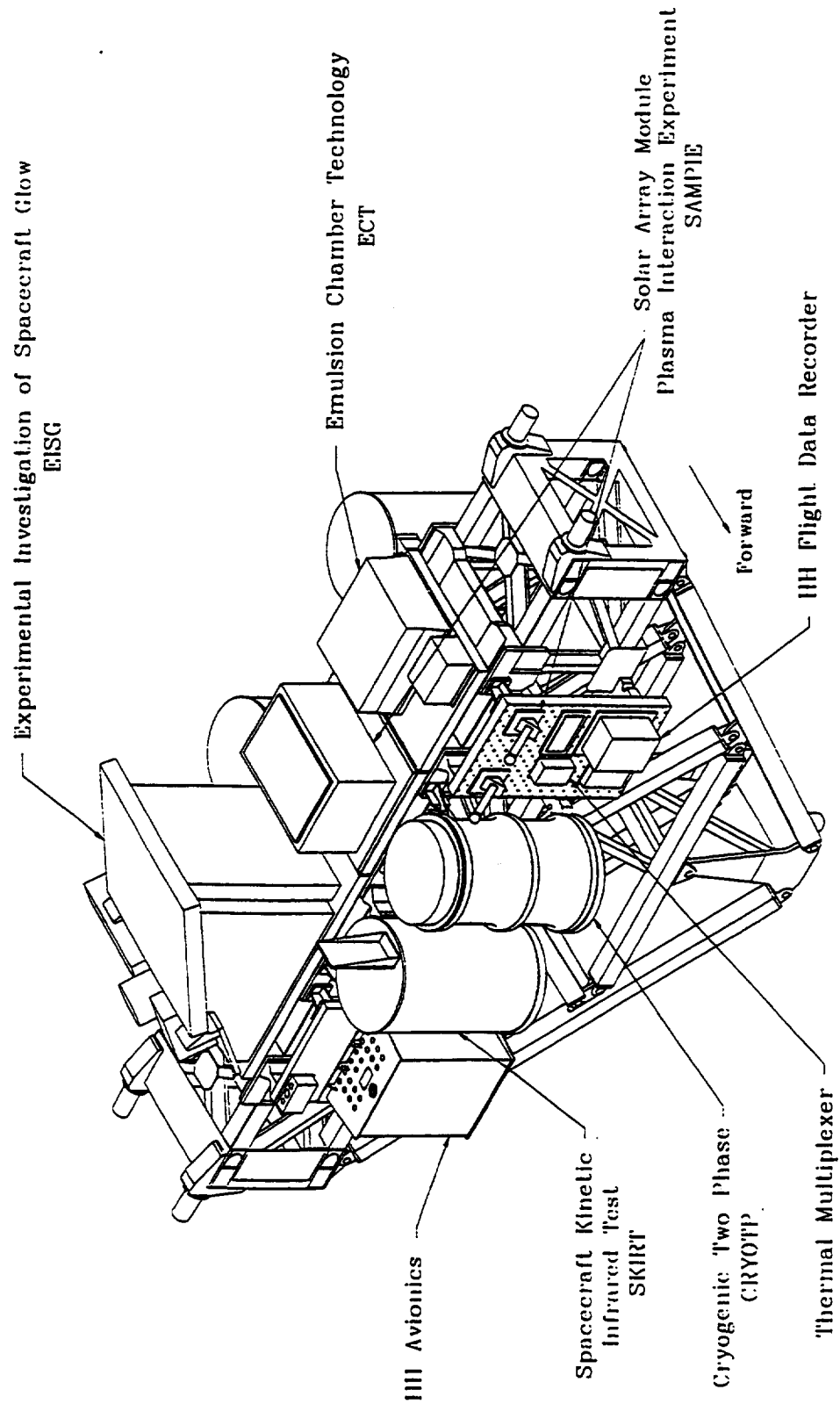


Figure 7

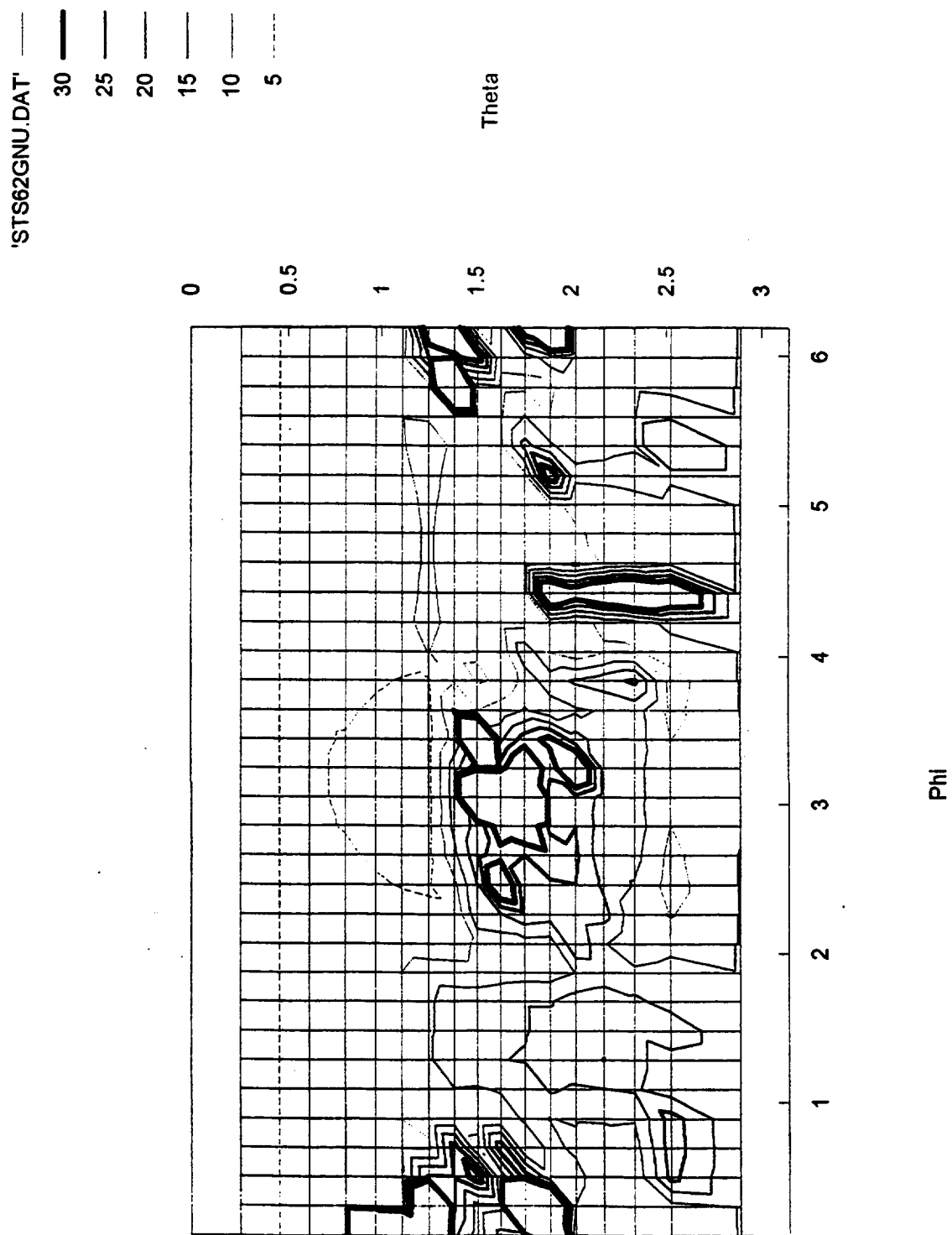


Figure 8

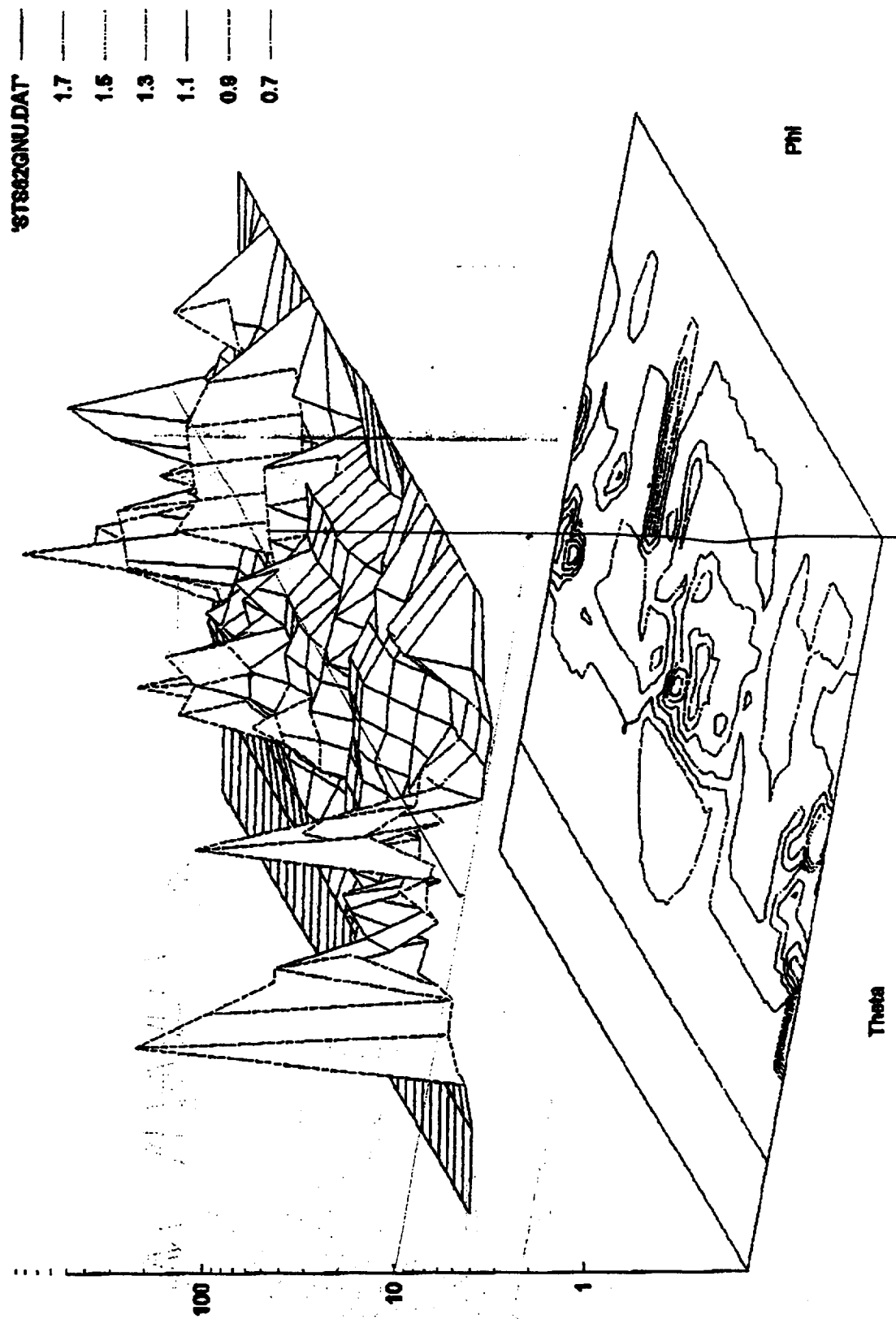


Figure 9

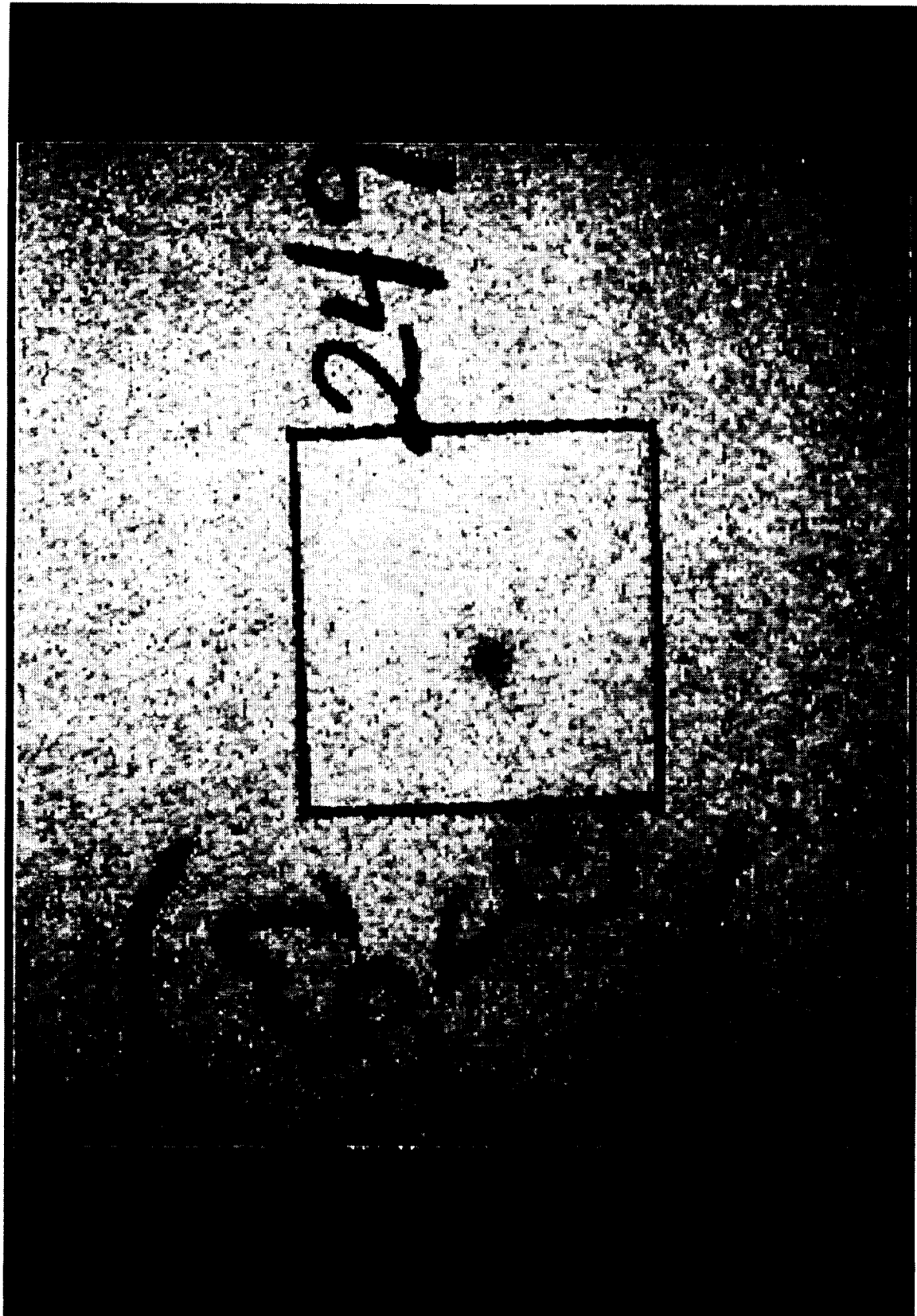
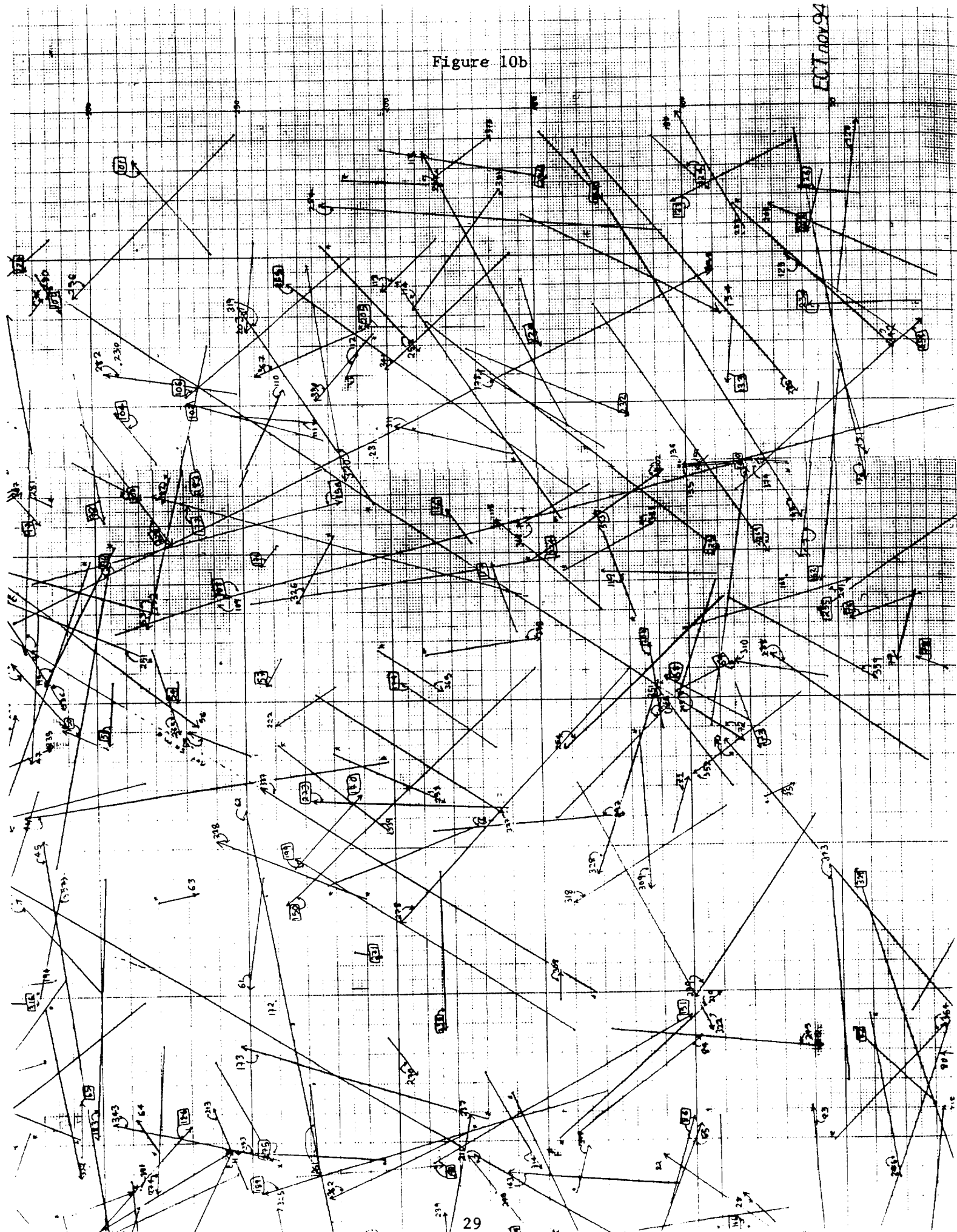


Figure 10a

Figure 10b

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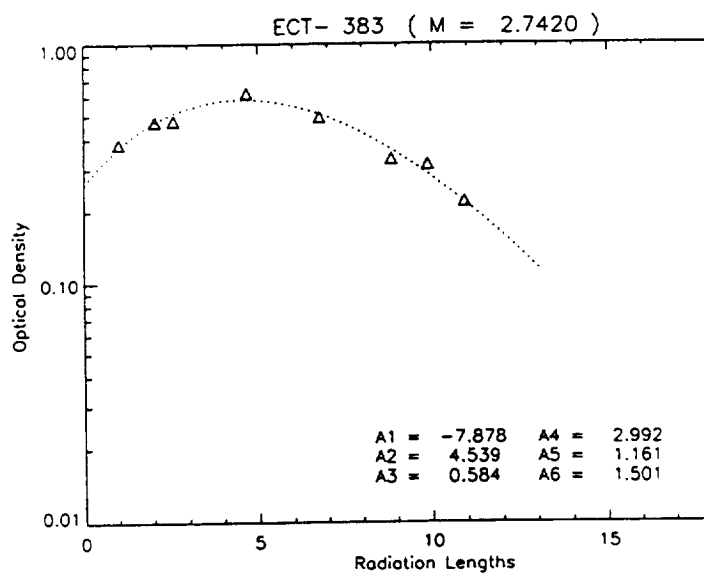
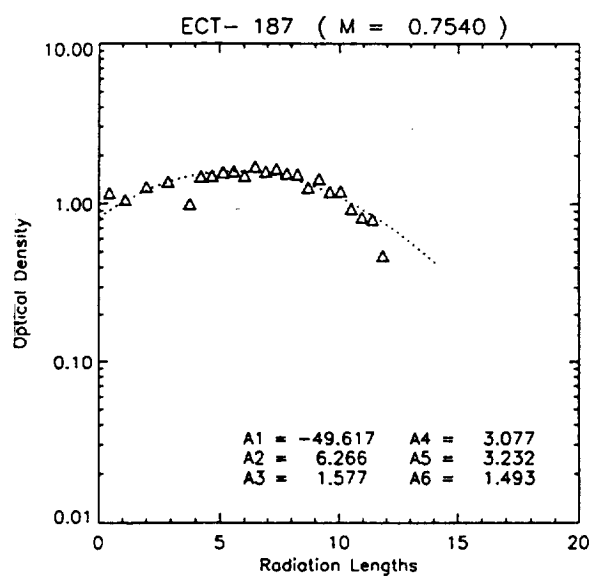
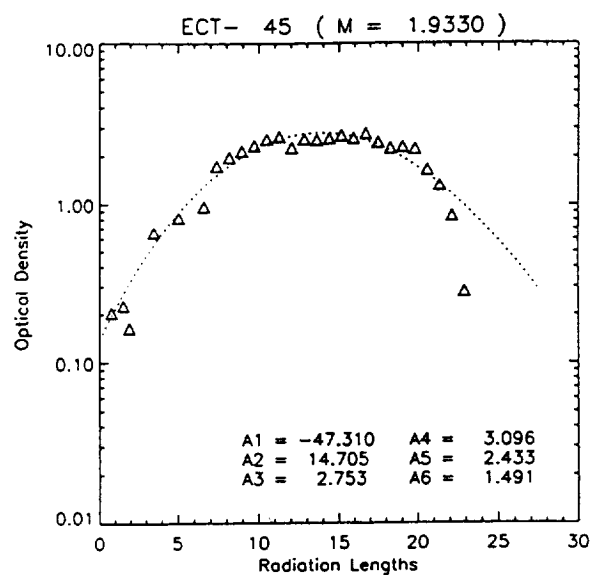
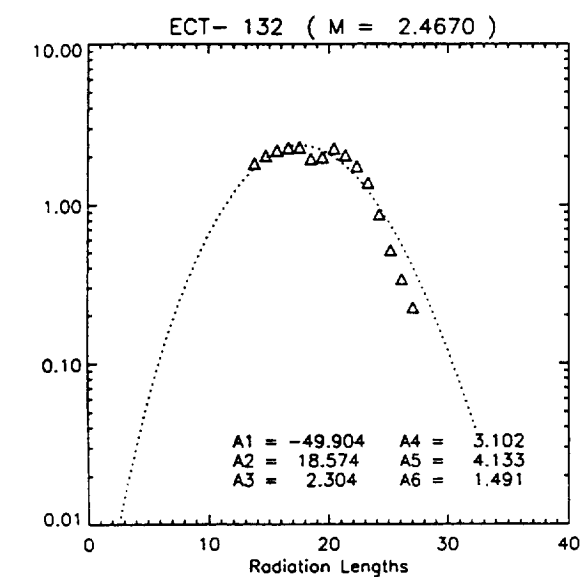


Figure 11

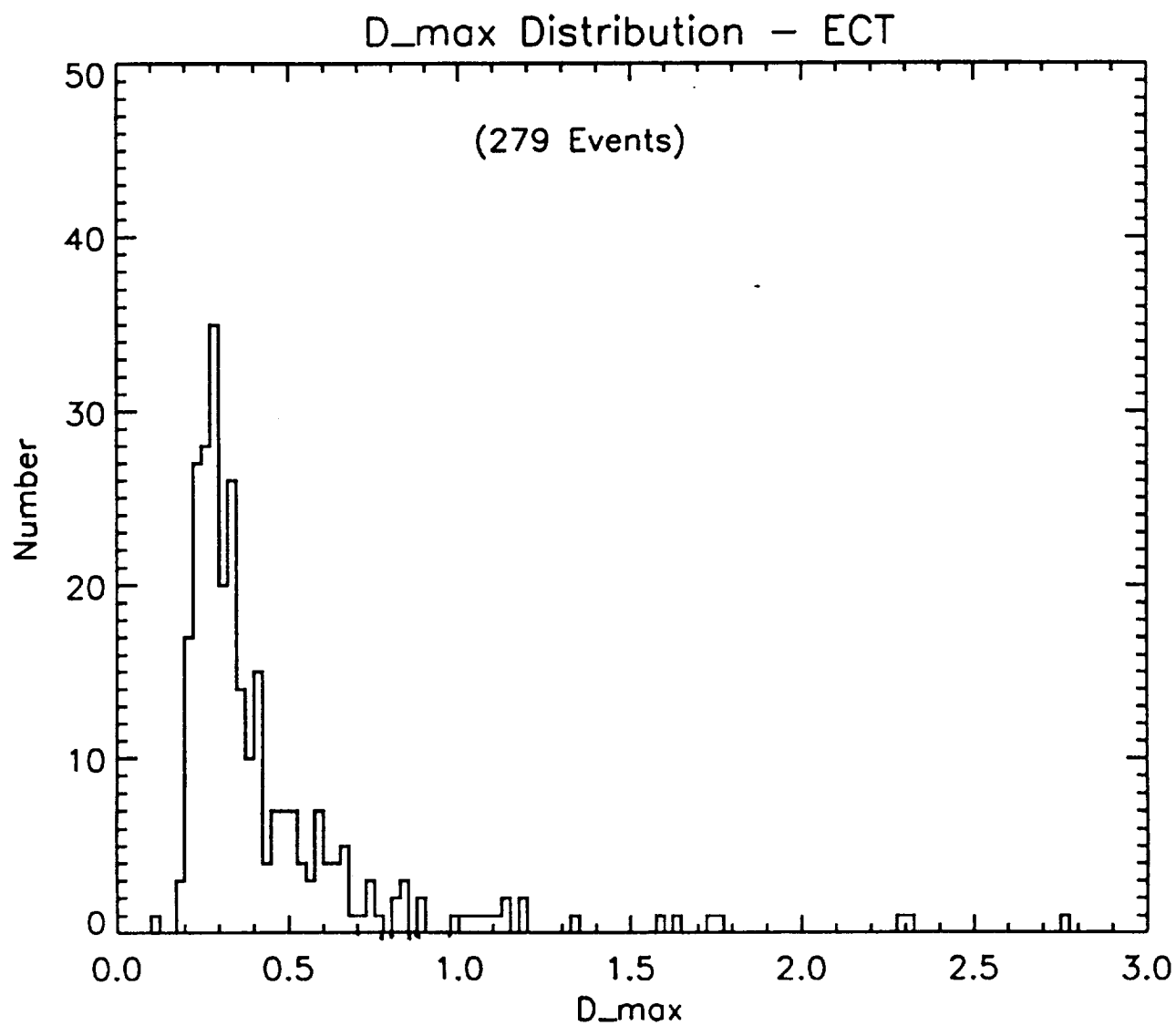


Figure 12a

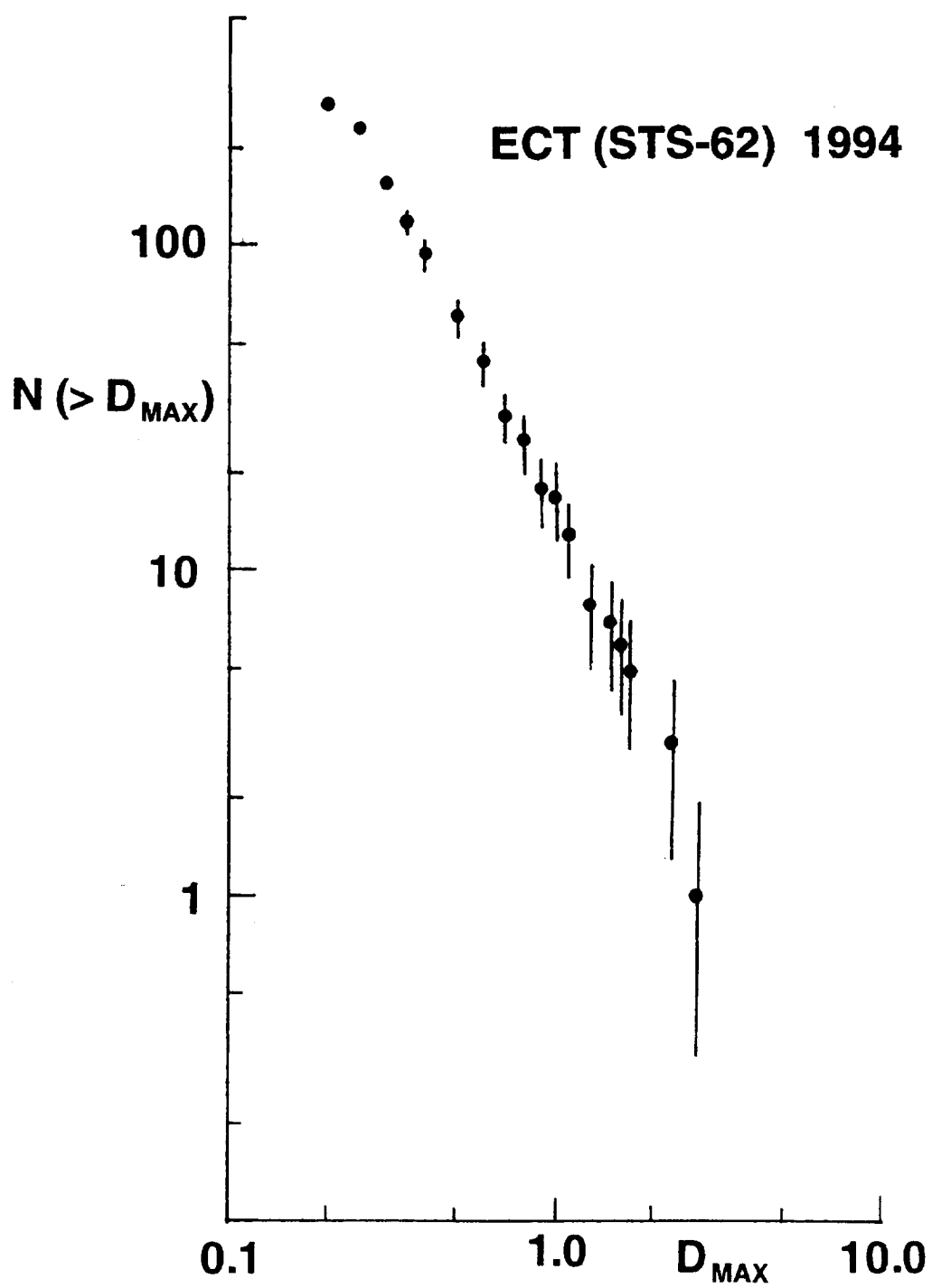


Figure 12b

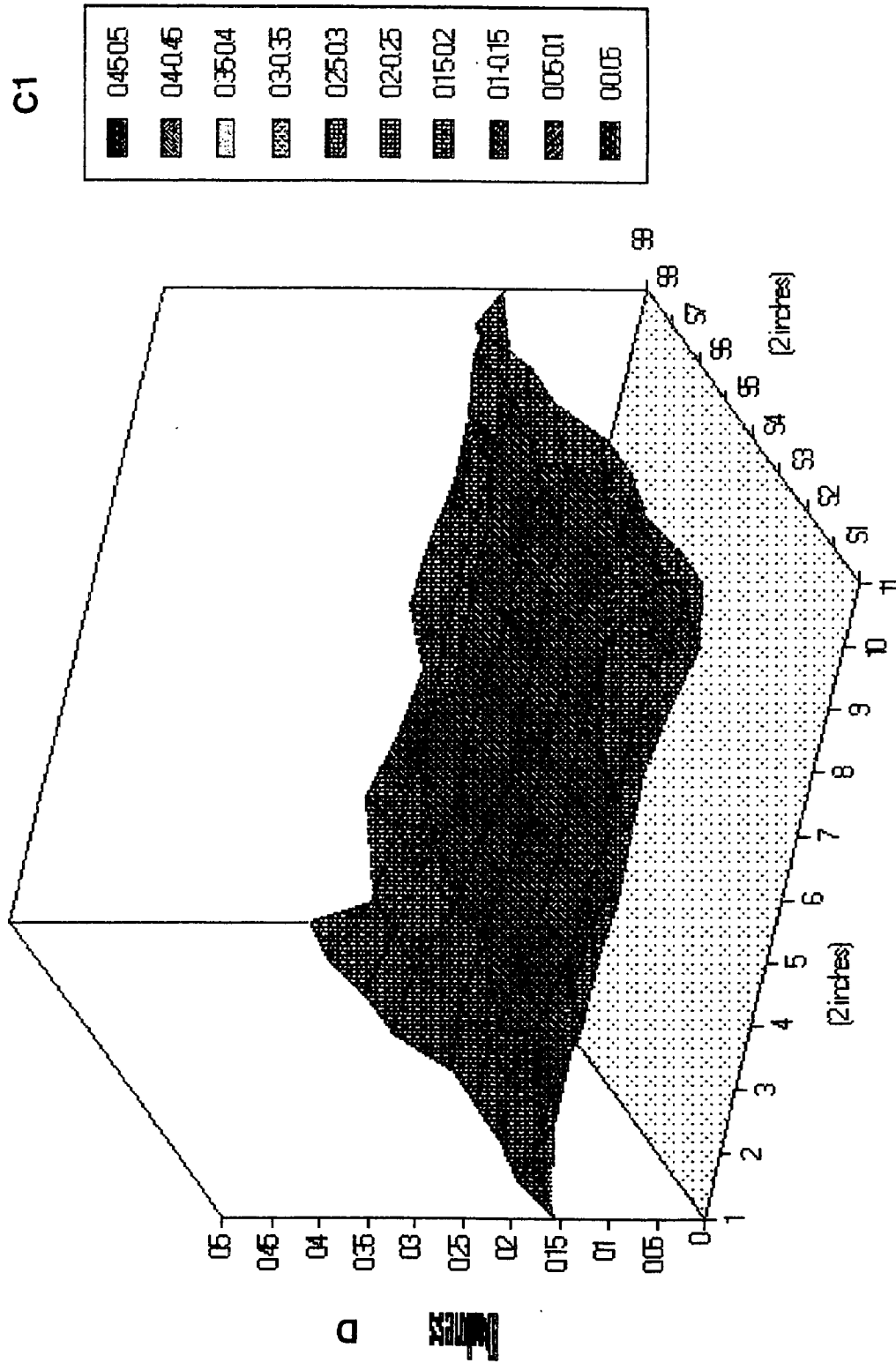


Figure 13-1

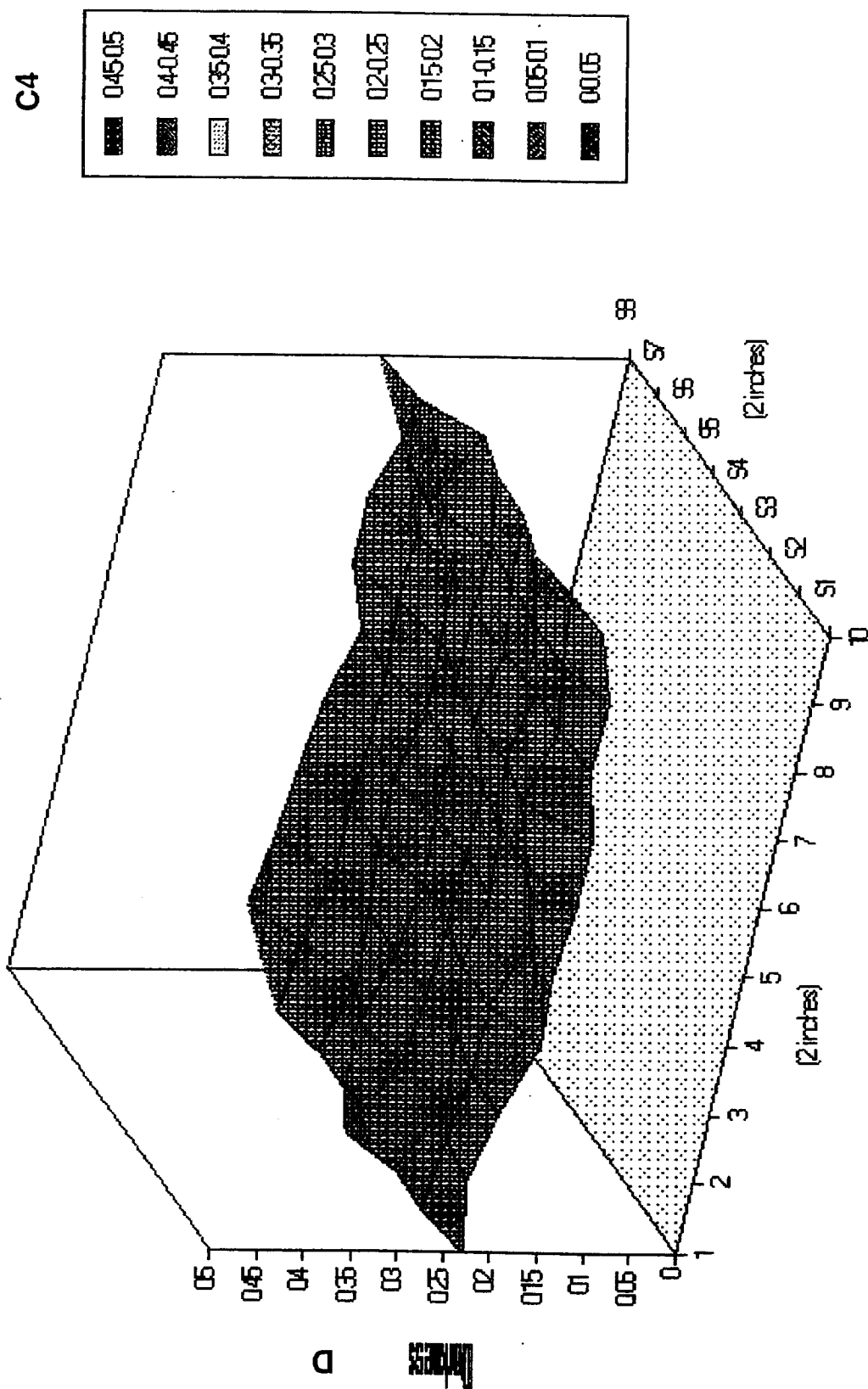


Figure 13-2

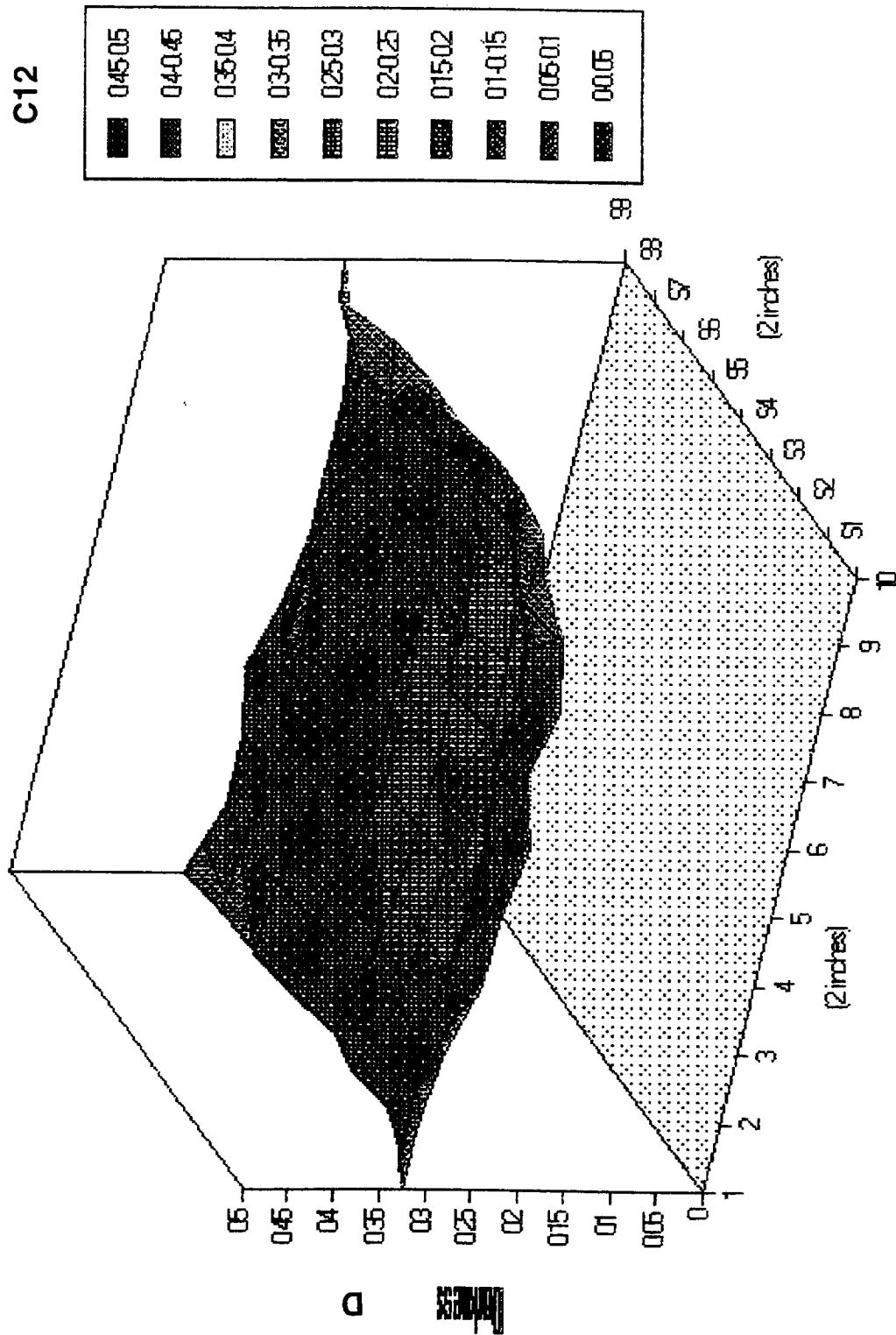


Figure 13-3

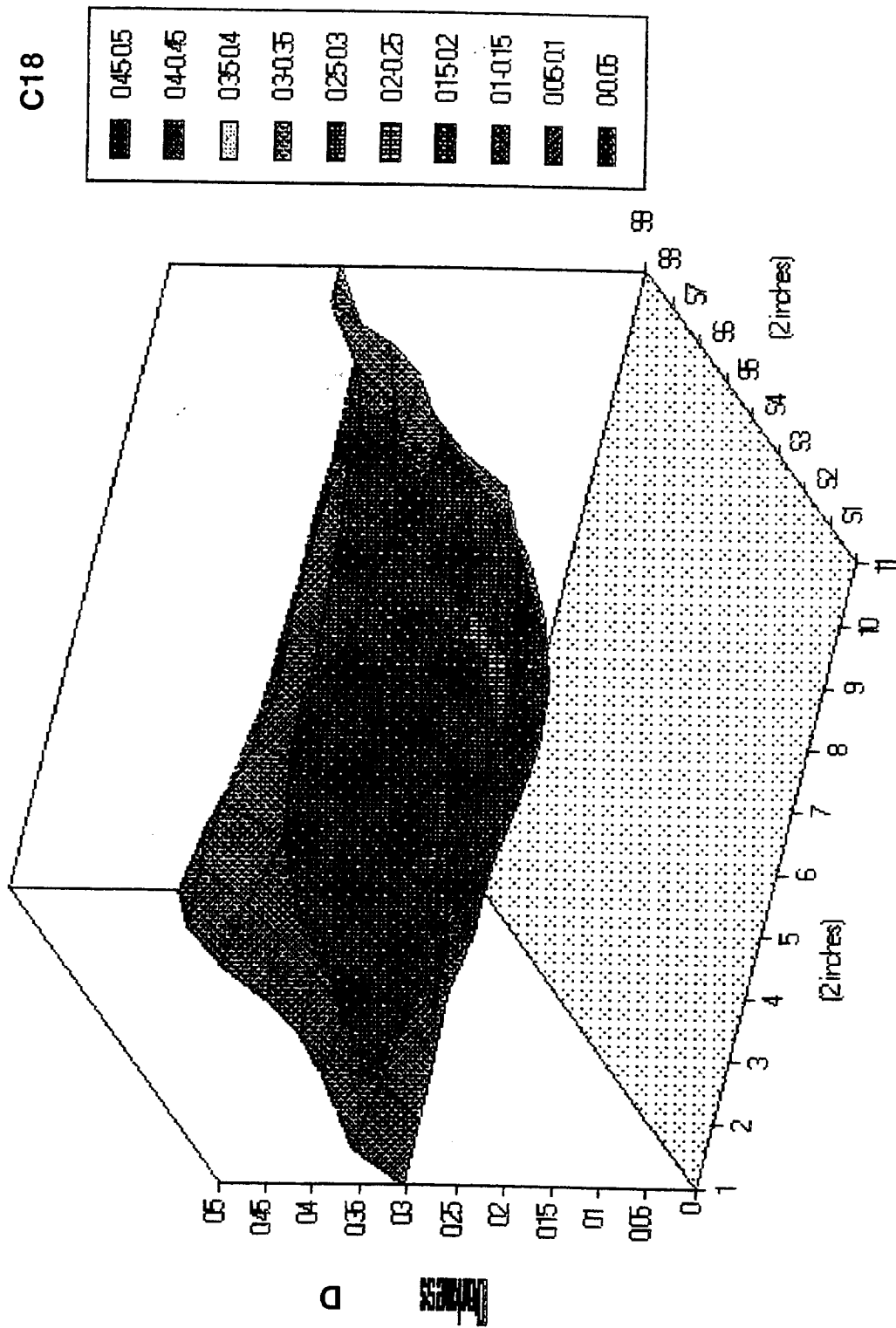


Figure 13-4

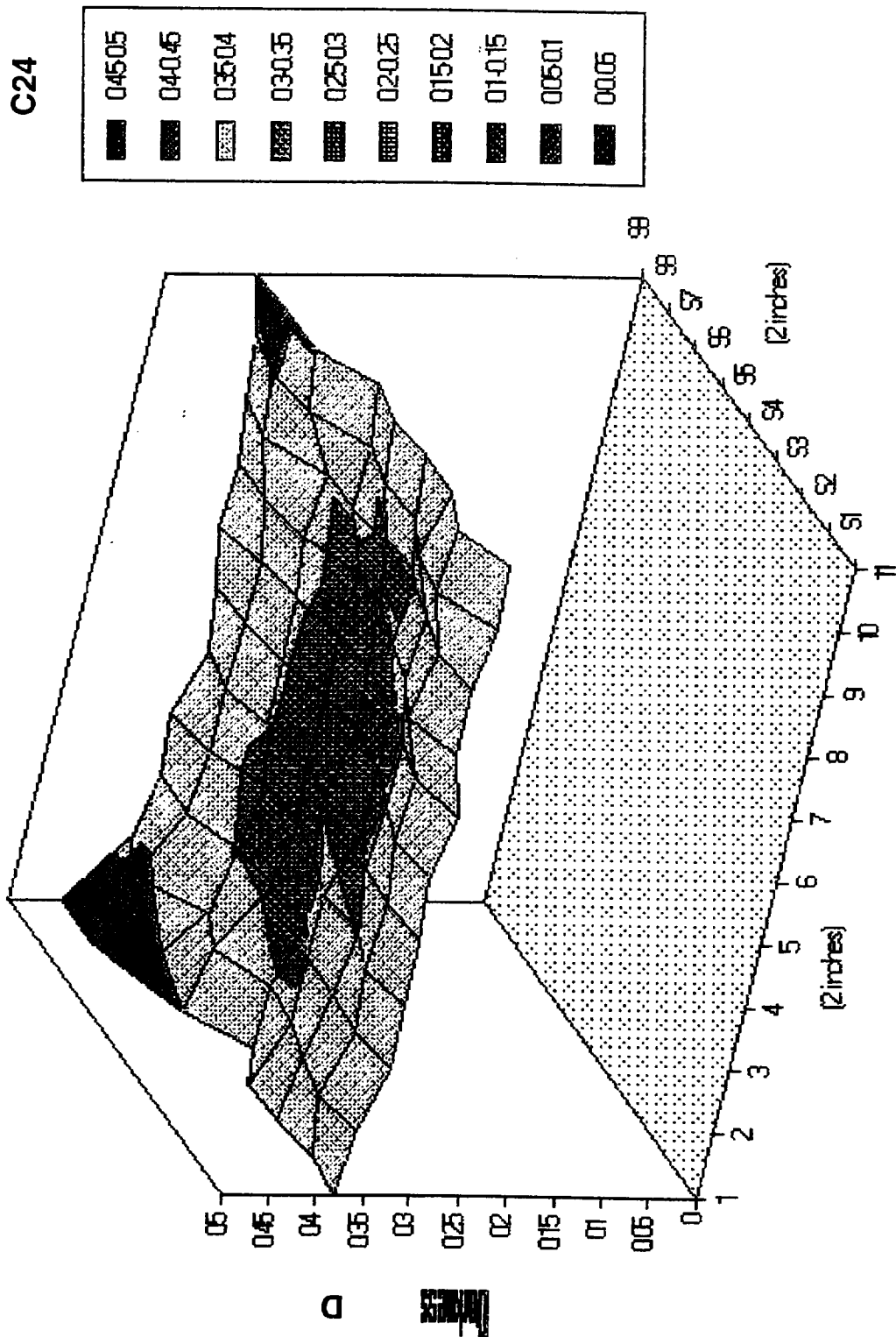


Figure 13-5

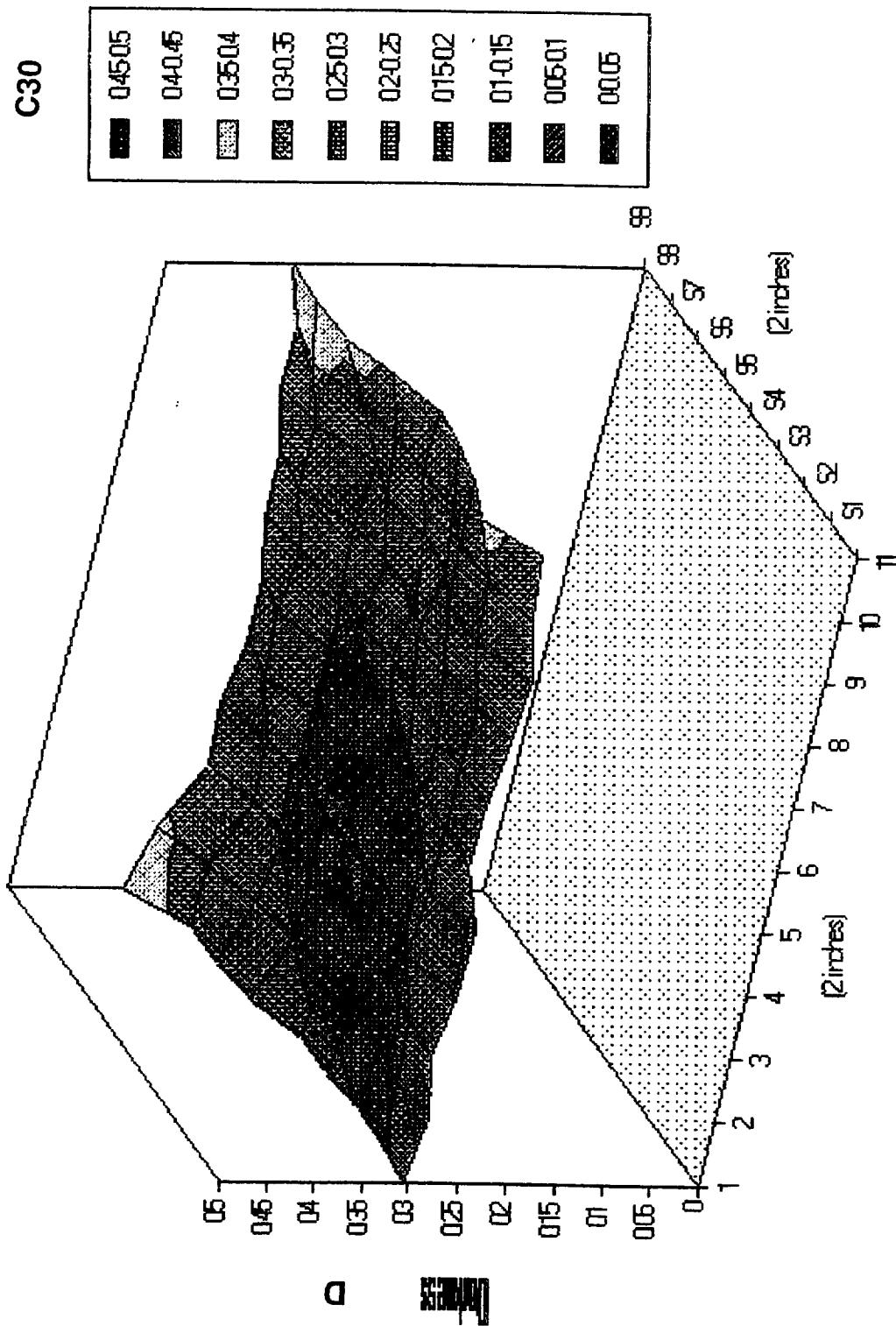


Figure 13-6

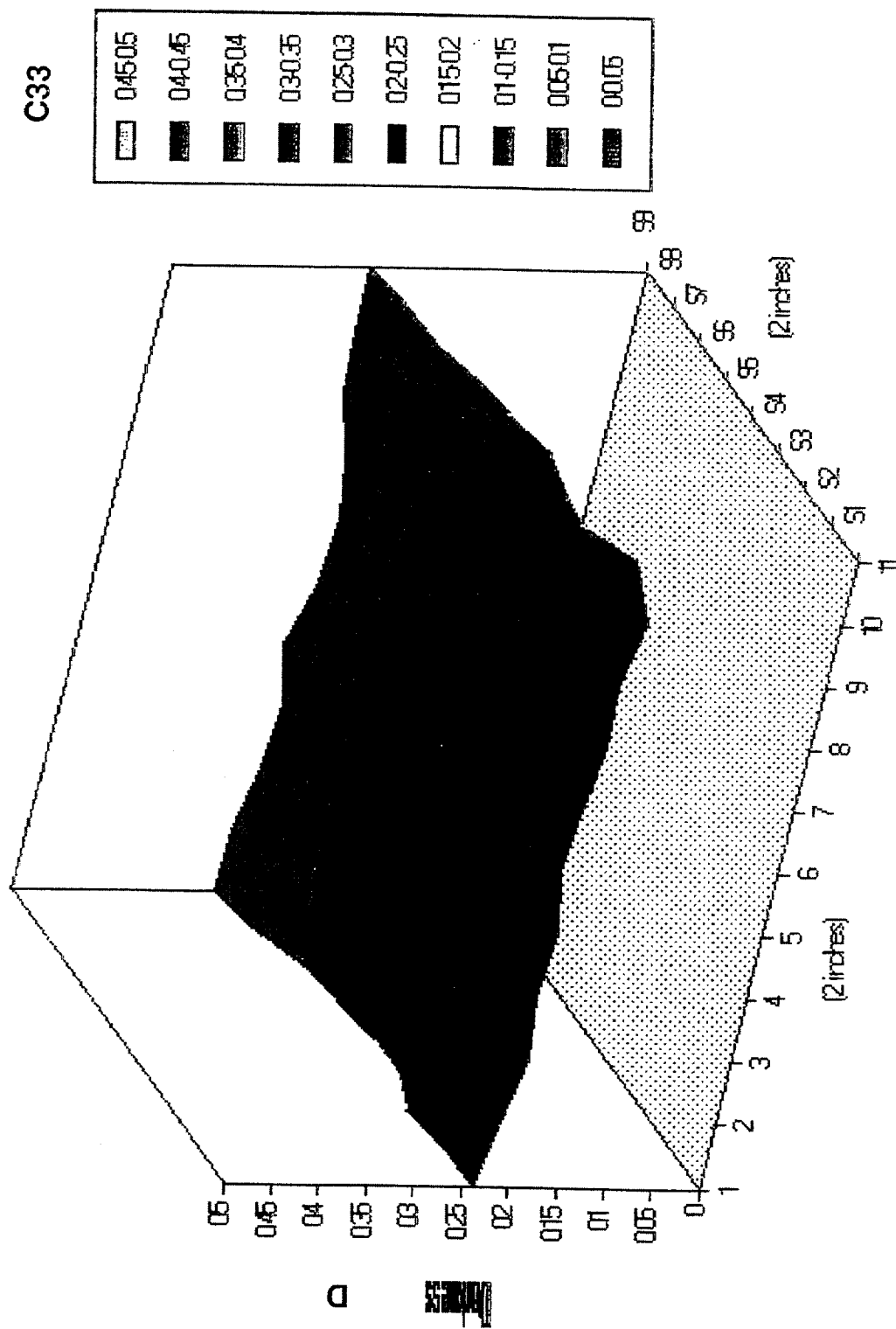


Figure 13-7

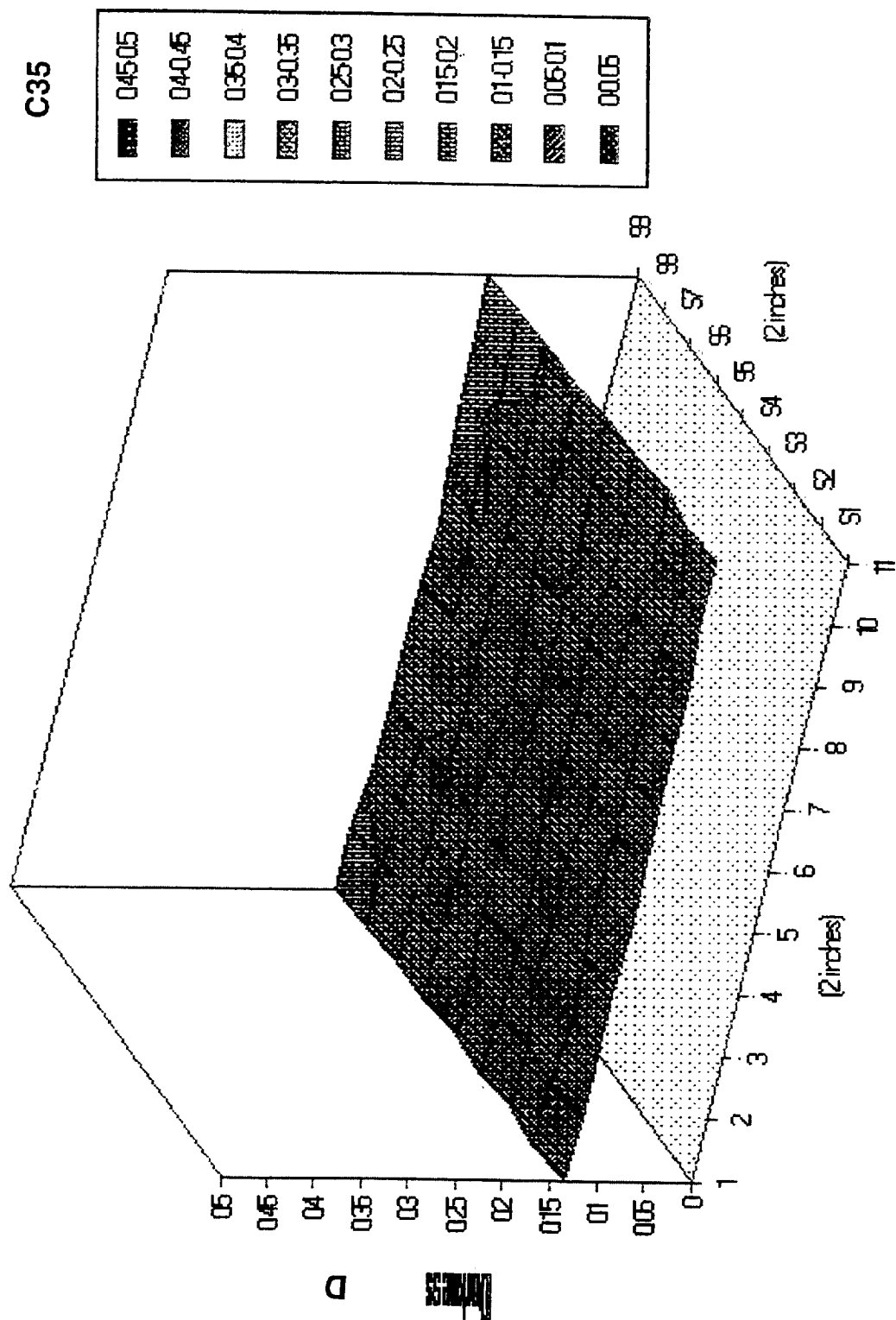
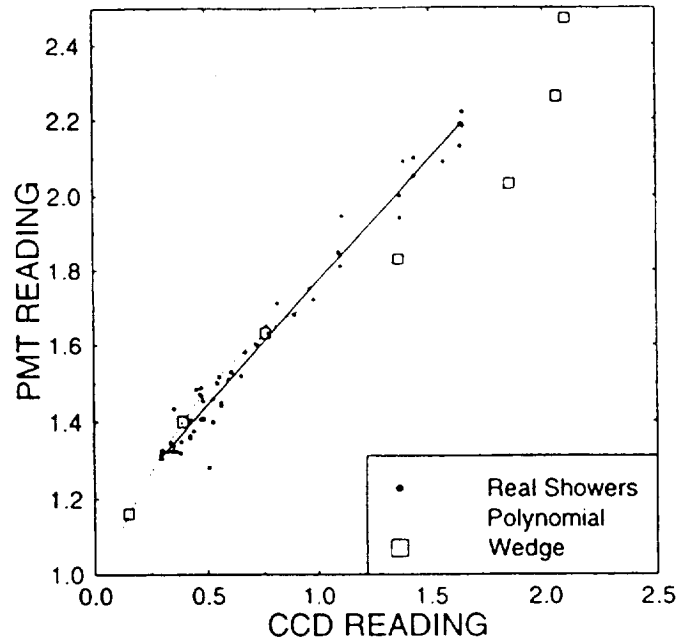
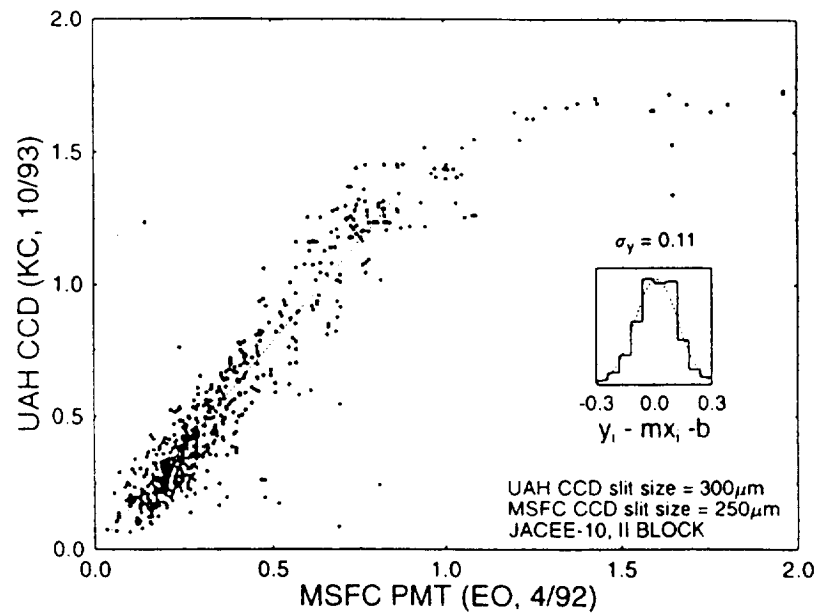


Figure 13-8



Raw CCD density calibration. Slit size is $250\mu\text{m}$ for both systems. Note the distinct difference between sterile (wedge) and dirty (real data) calibration.



Net CCD density calibration. Several hundred real shower spots are used. Measurements fit well within a line of $D_{CCD} = 1.55 D_{PMT}$ for $D_{PMT} < 1.0$

Figs. 14a & 14b

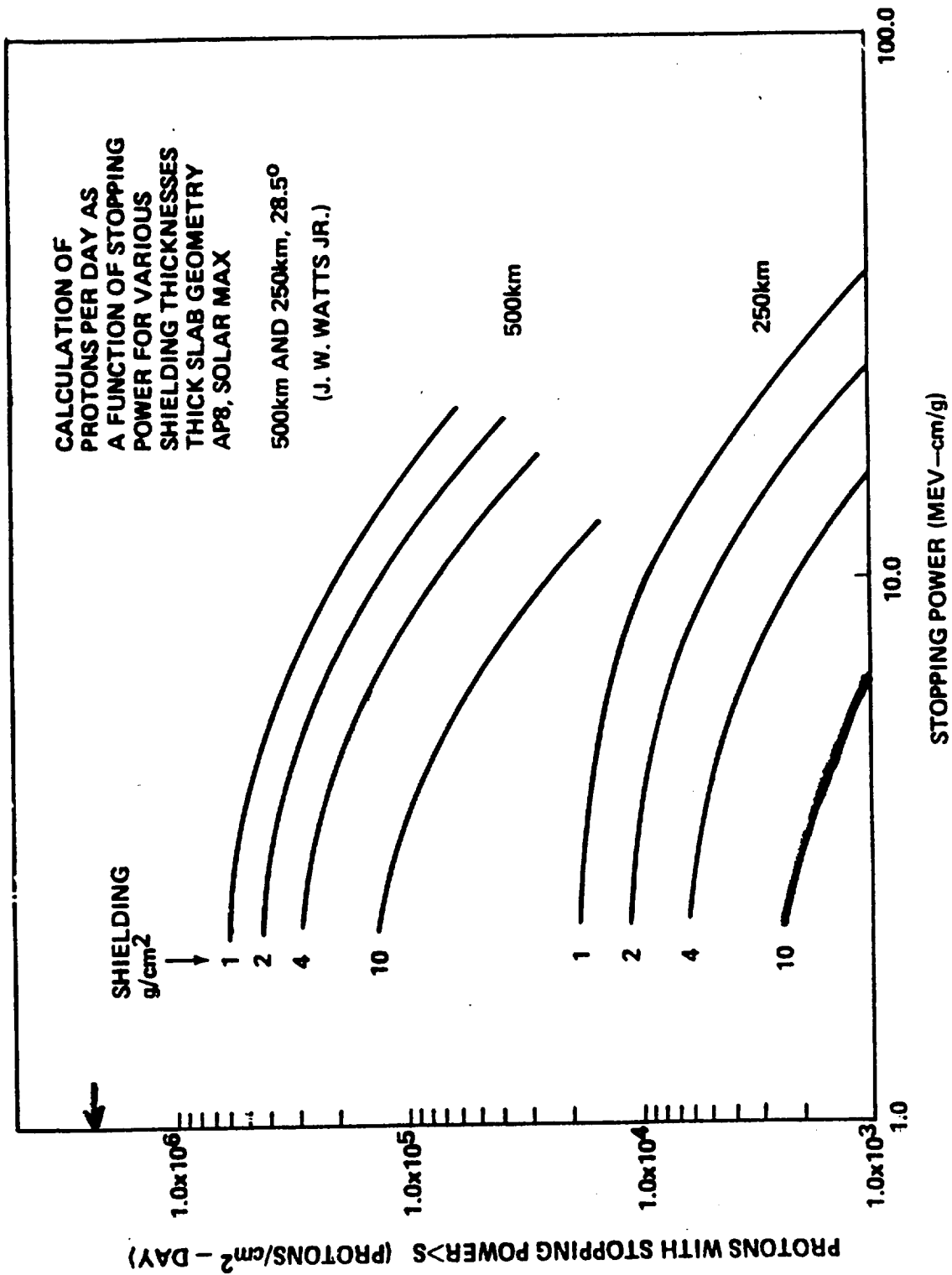


Figure 14c

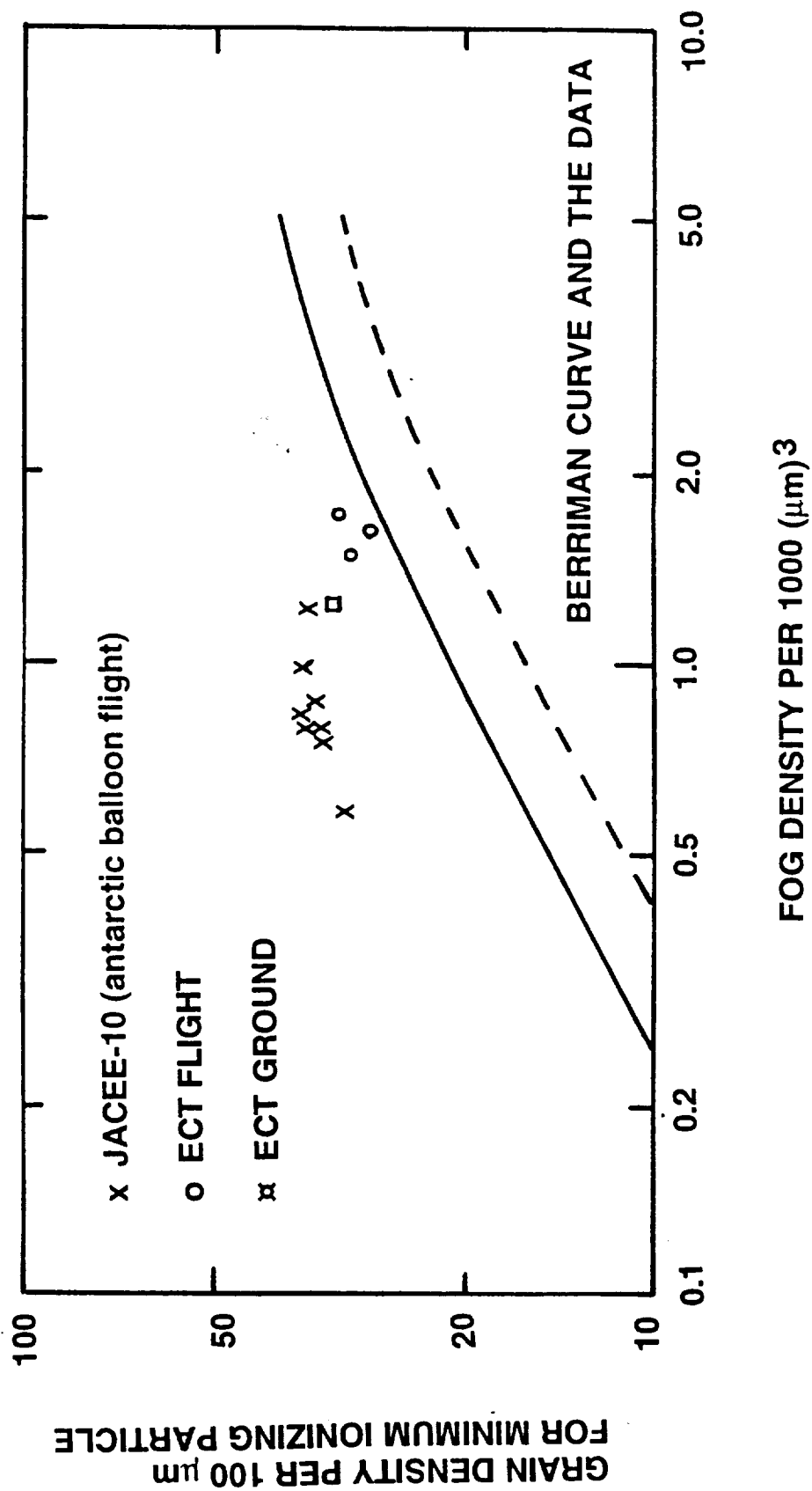


Figure 15

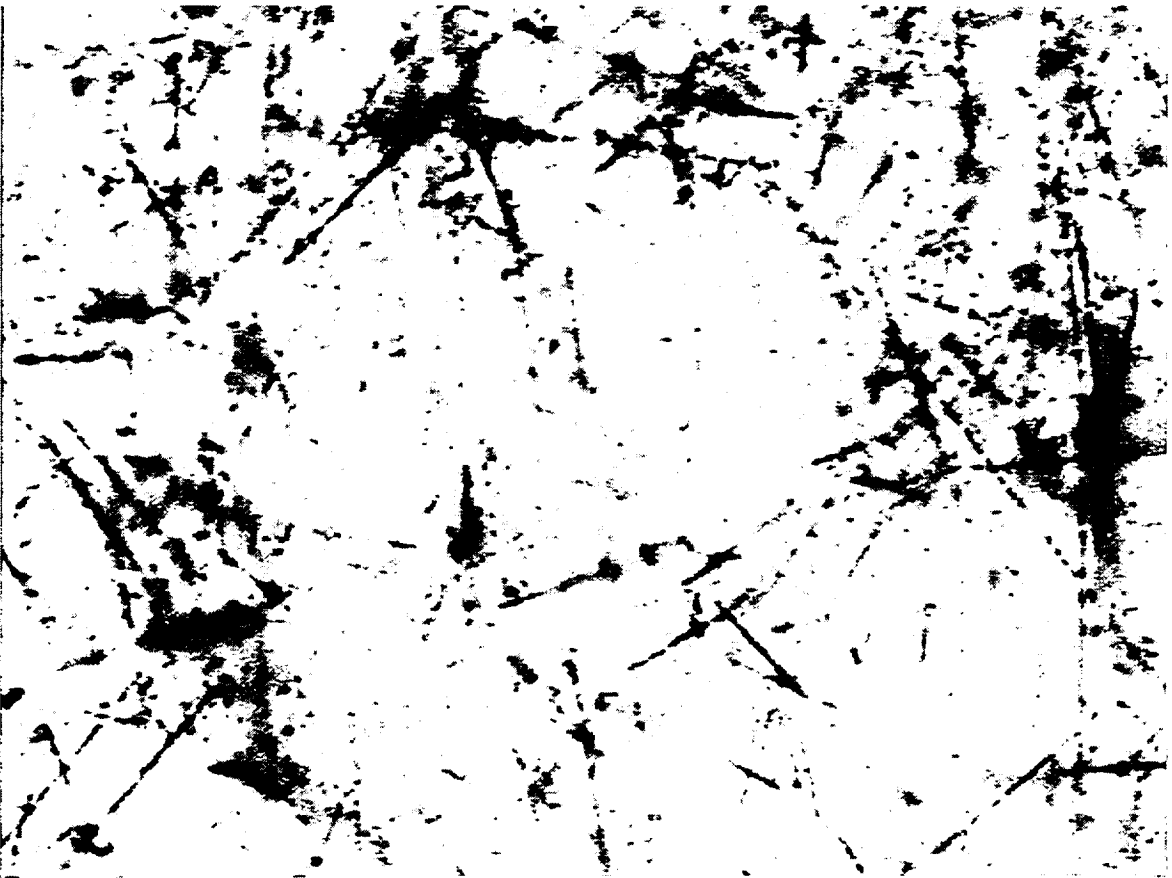
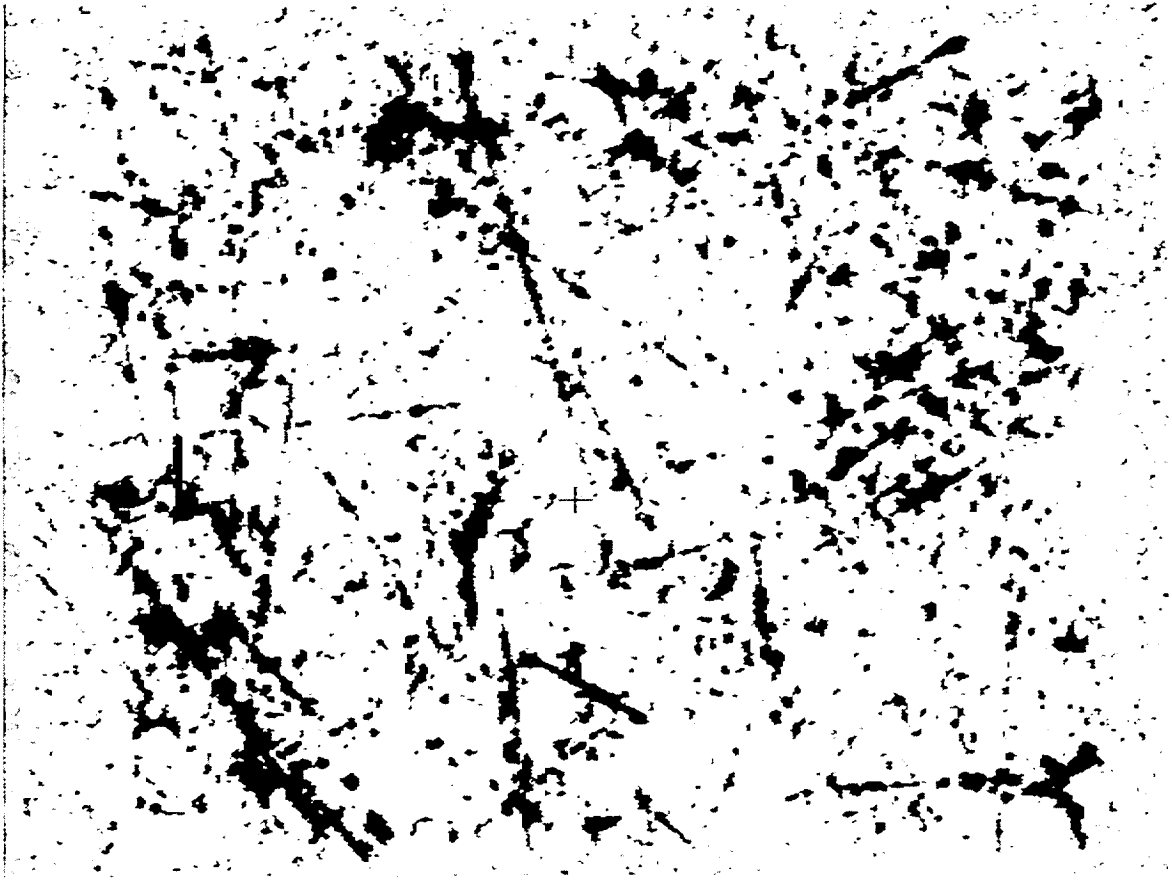


Figure 16

TempMentor Data for 1/6/94 to 3/27/94

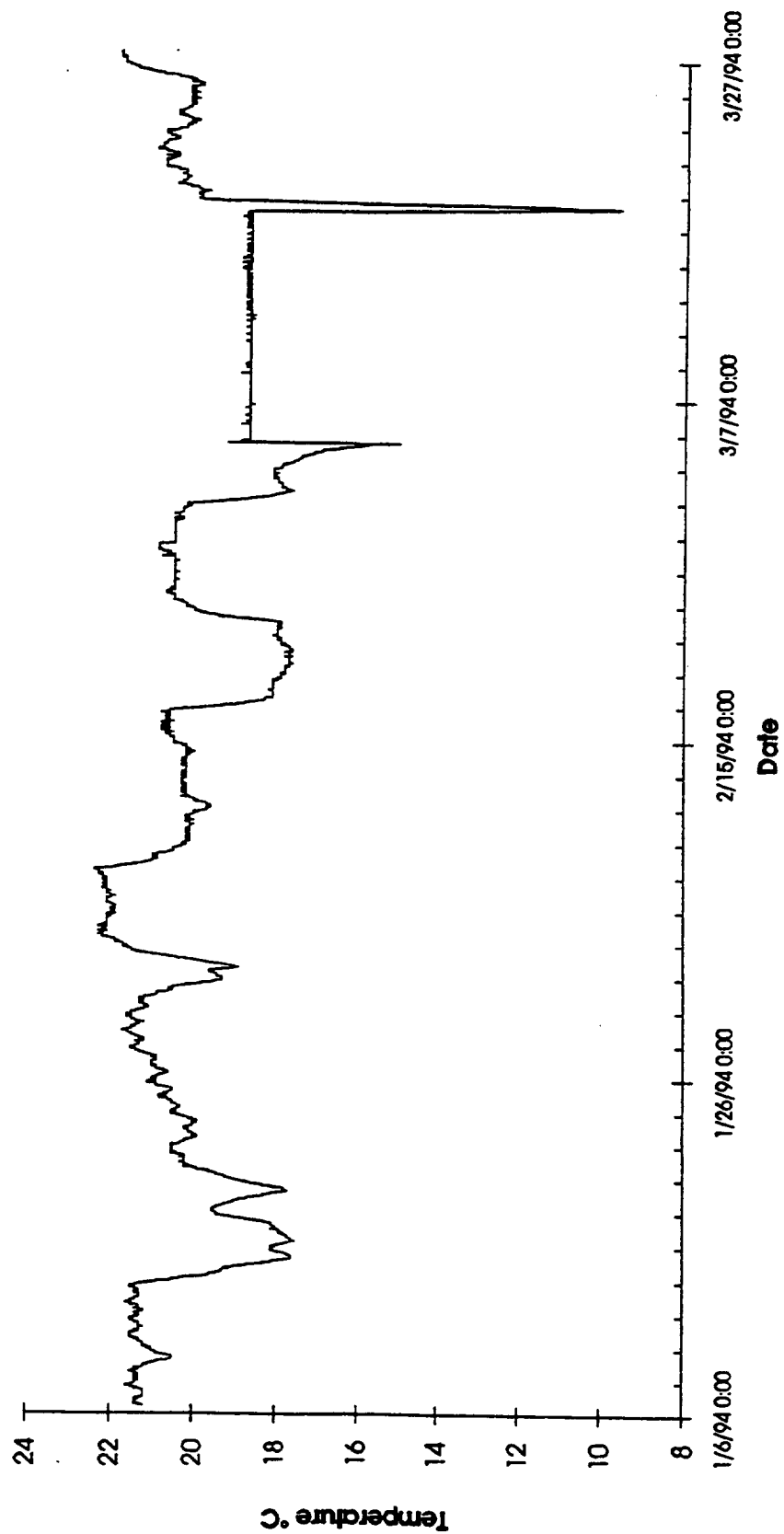
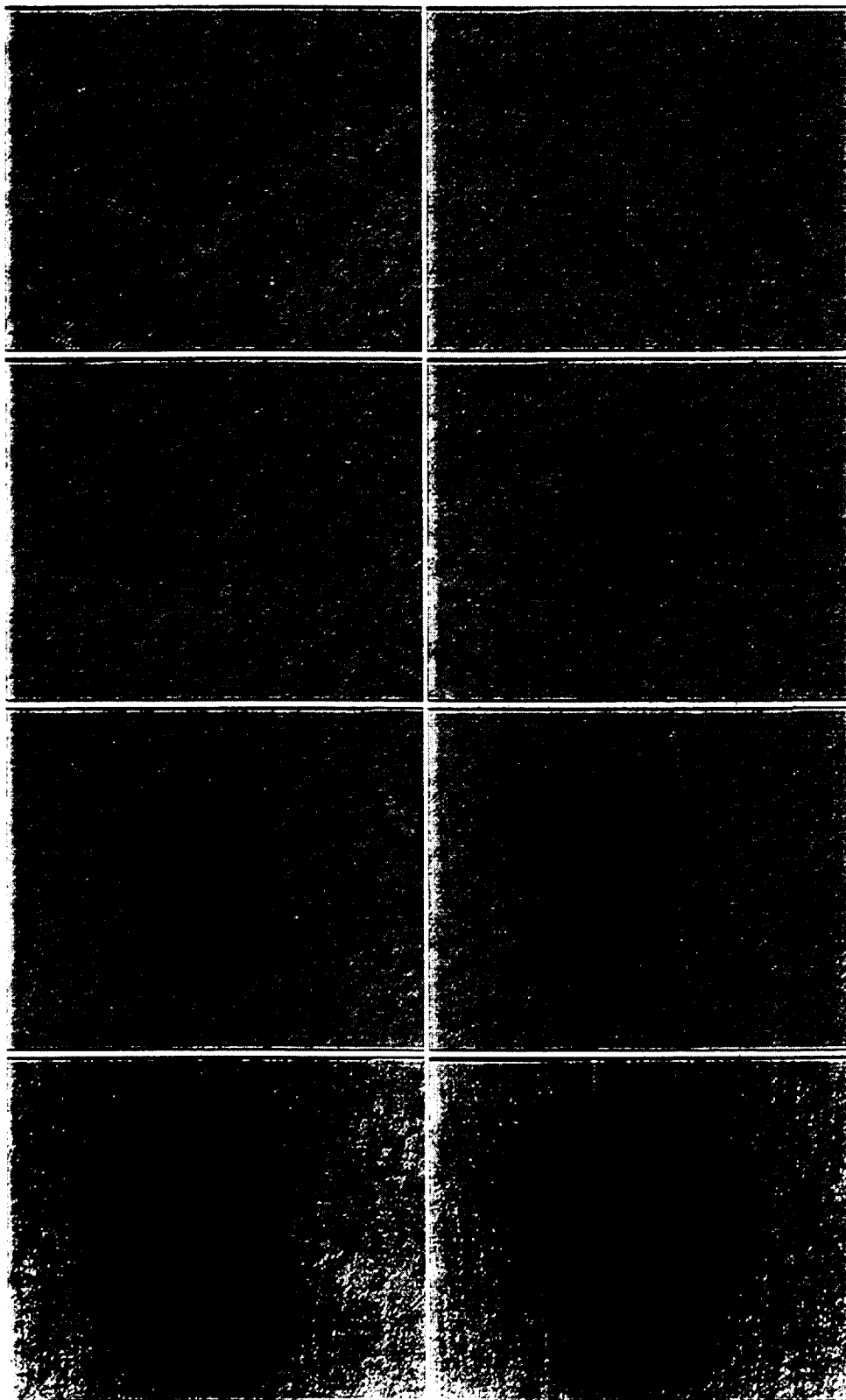
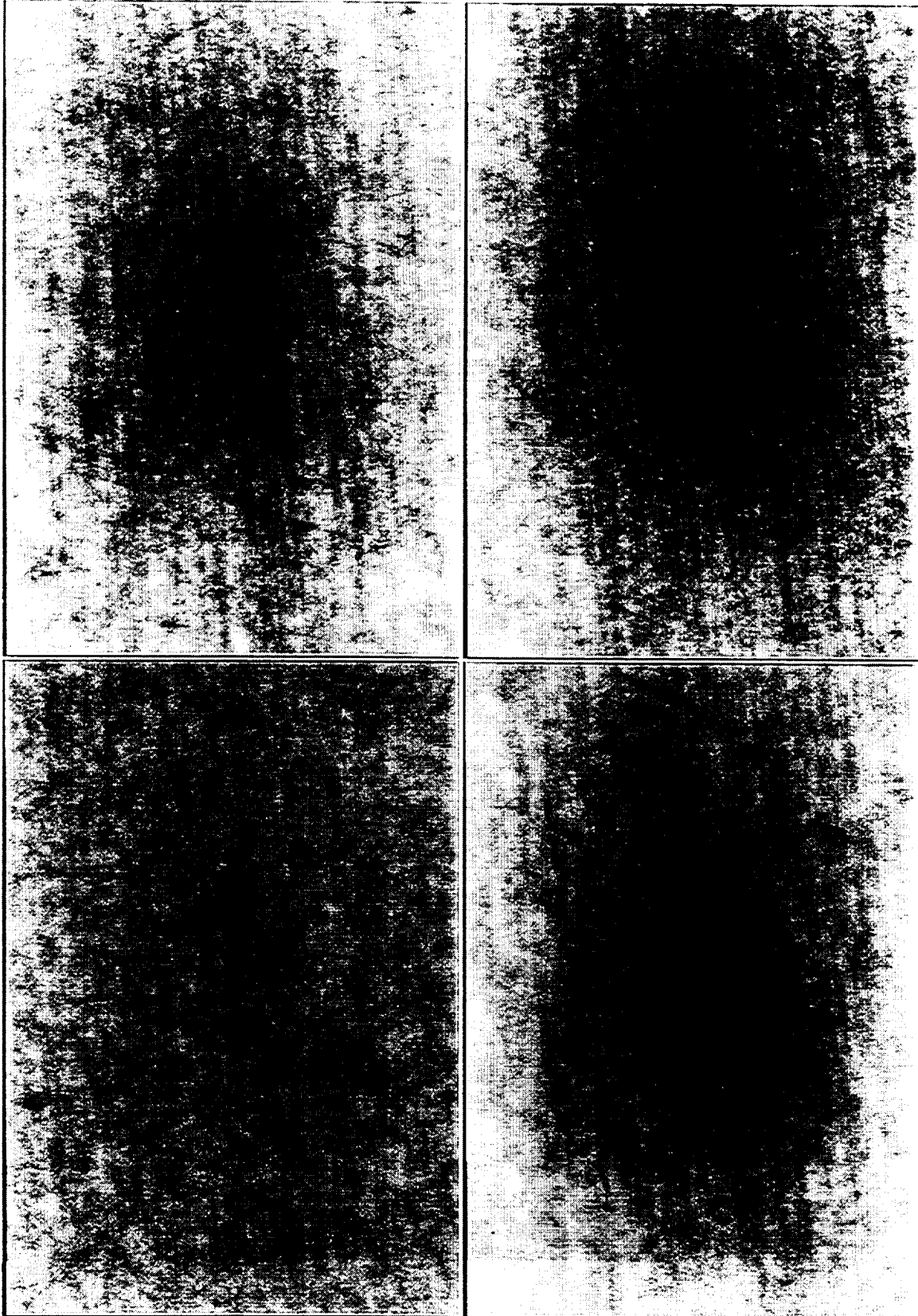


Figure 17



ECT FLIGHT EMULSIONS P01/03/21/69/C01/05/11/13

Figure 18



ICI 11 IGH C12/11/21/25

Figure 19



ECT FLIGHT EMULSIONS

Figure 20



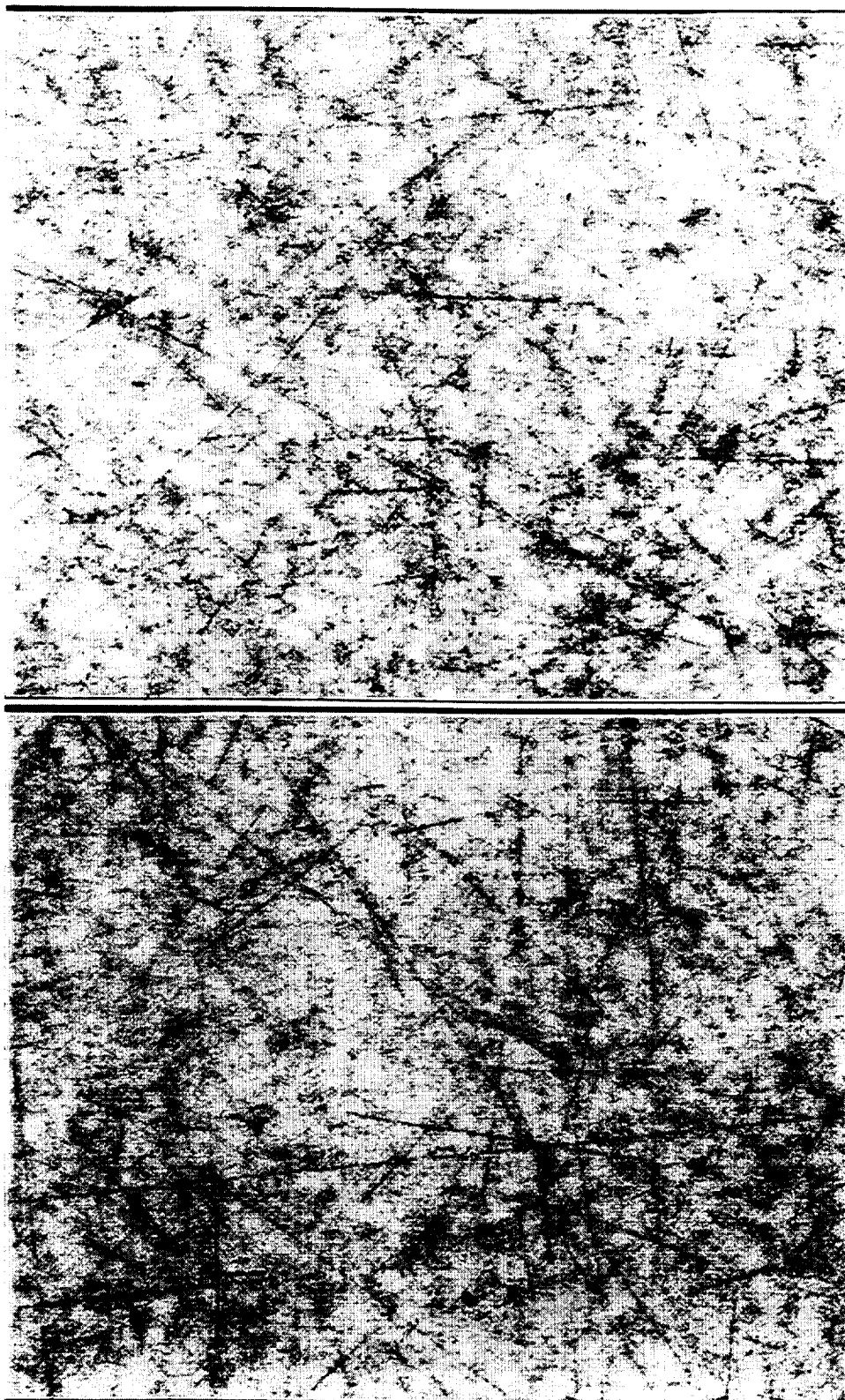
ECT FLIGHT EMULSIONS

Figure 21



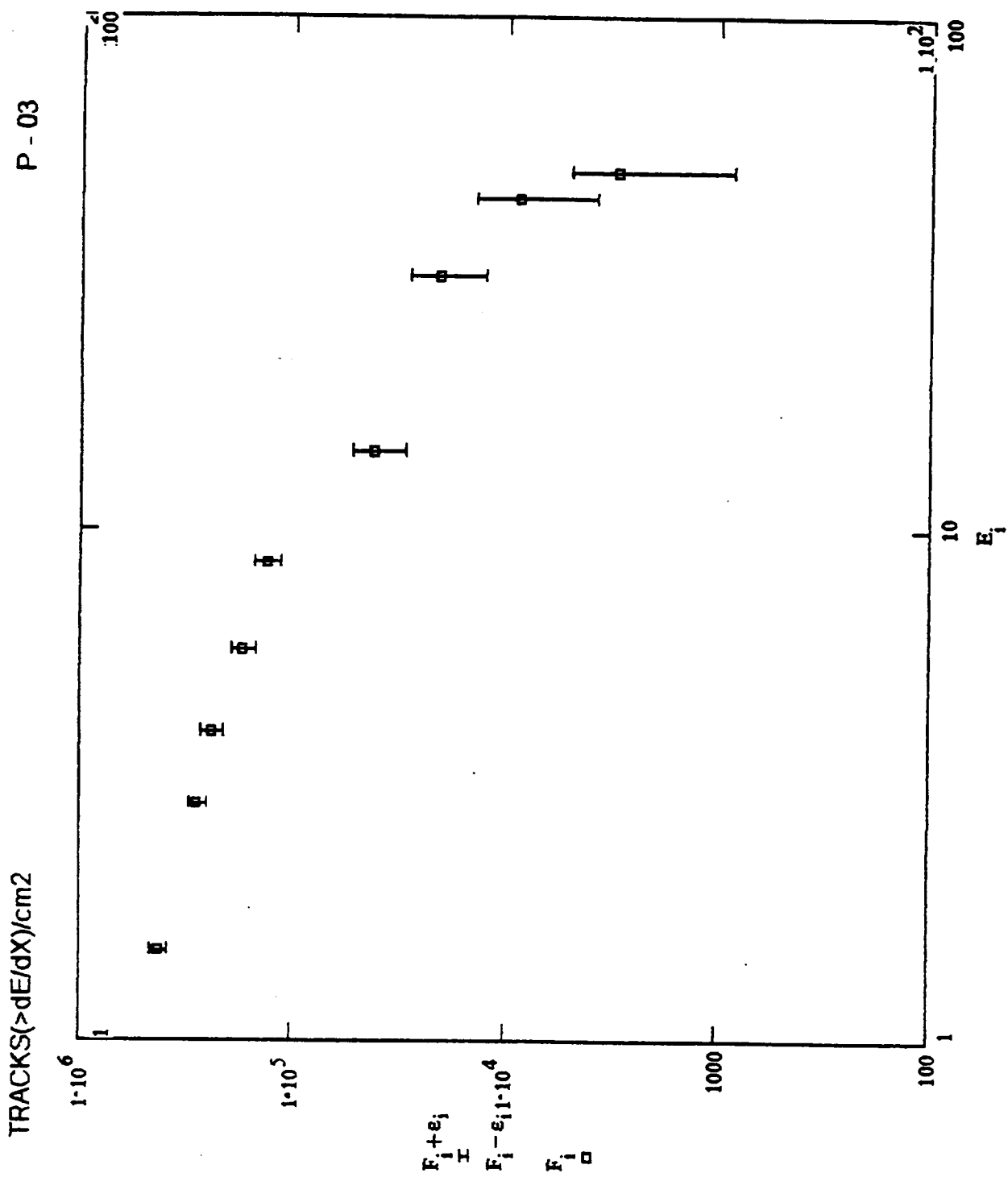
ECT FLIGHT EMULSIONS

Figure 22



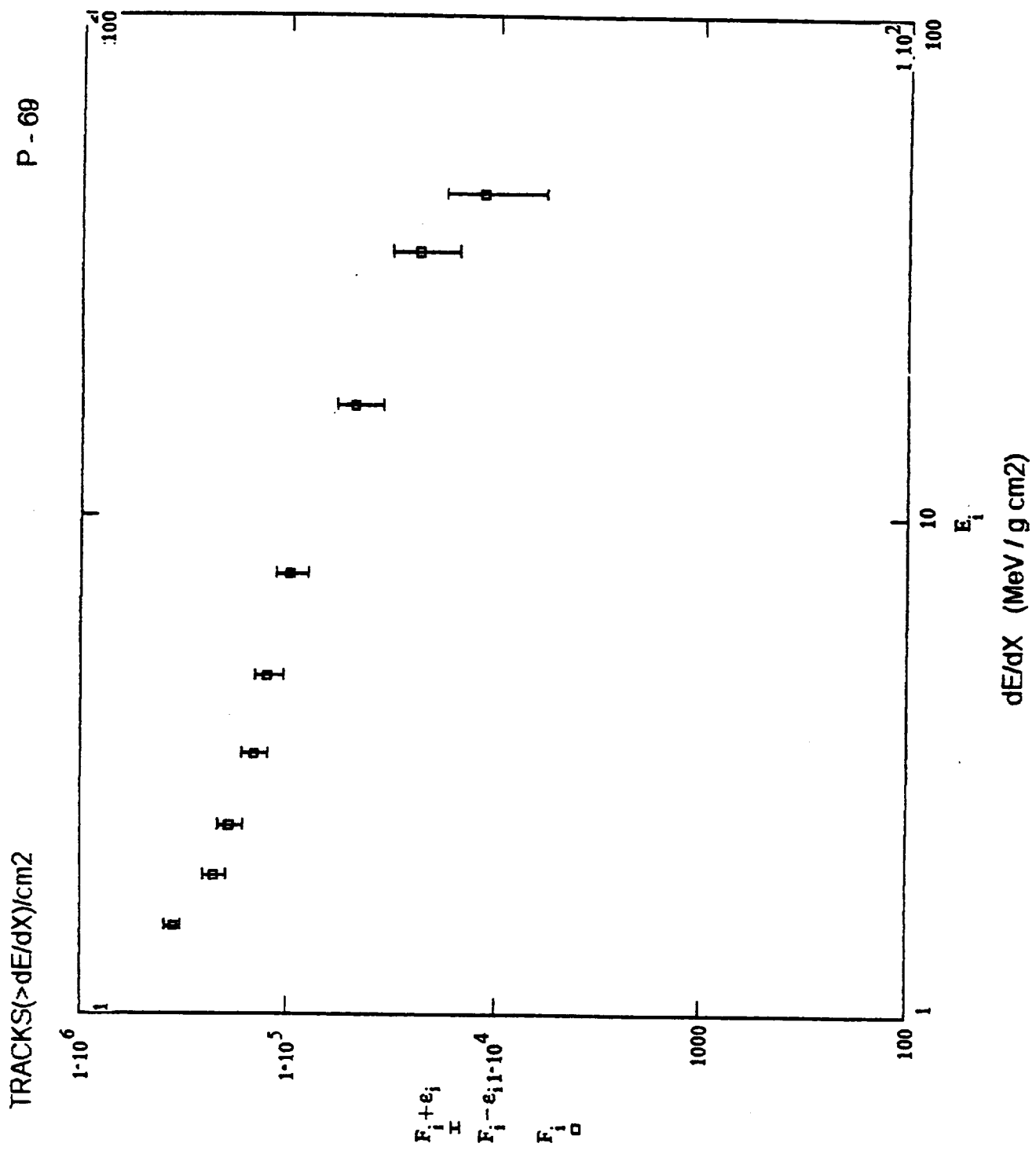
ECT FLIGHT (TOP) LDBF (BOTTOM)

Figure 23



Integral Linear Energy Transfer Spectrum
in the emulsion plate located 1.4 g/cm2
from the top of ECT chamber

Figure 24a



Integral Linear Energy Transfer Spectrum
in the emulsion plate located 16.7 g/cm2
from the top of ECT chamber

Figure 24-b

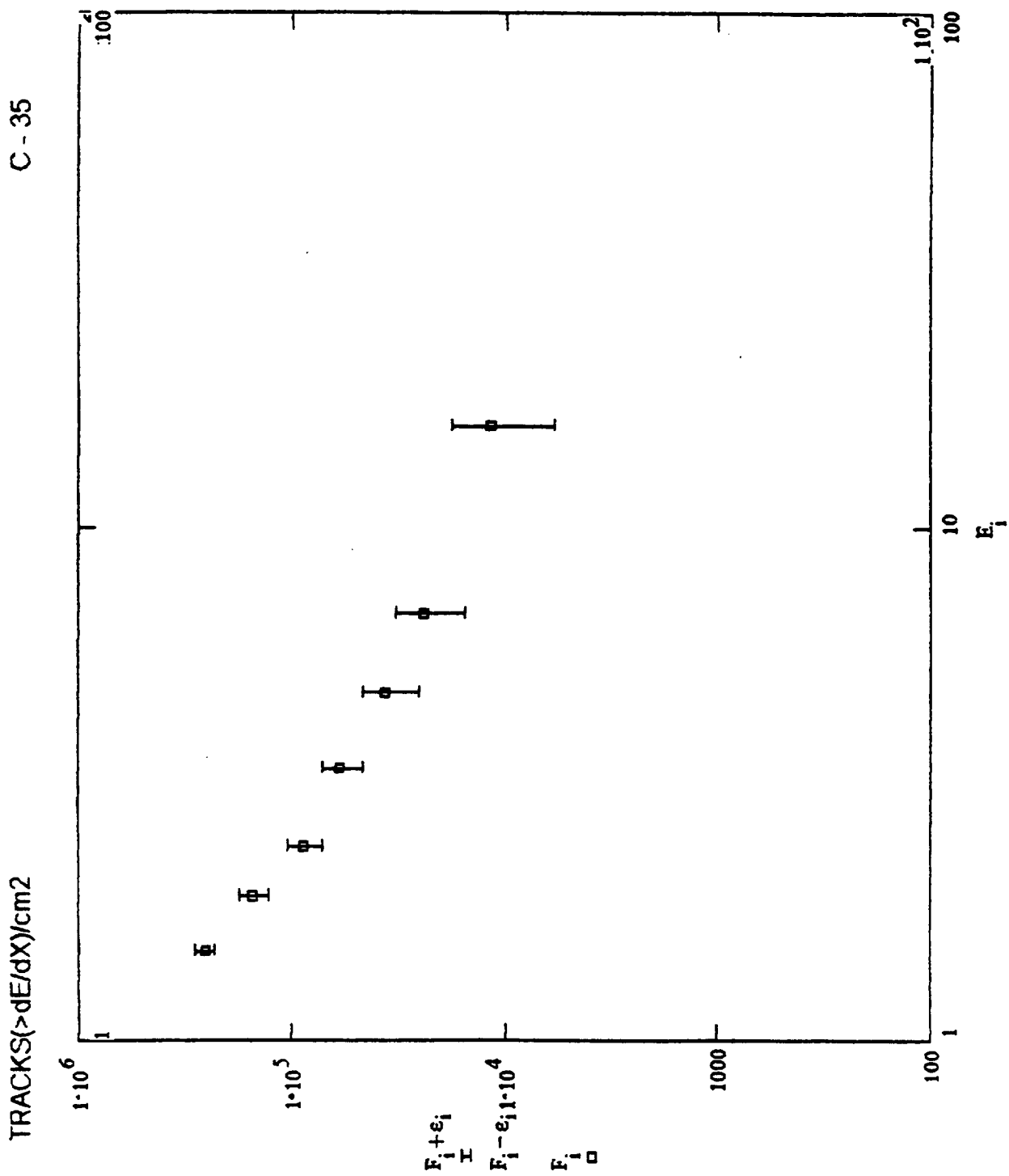
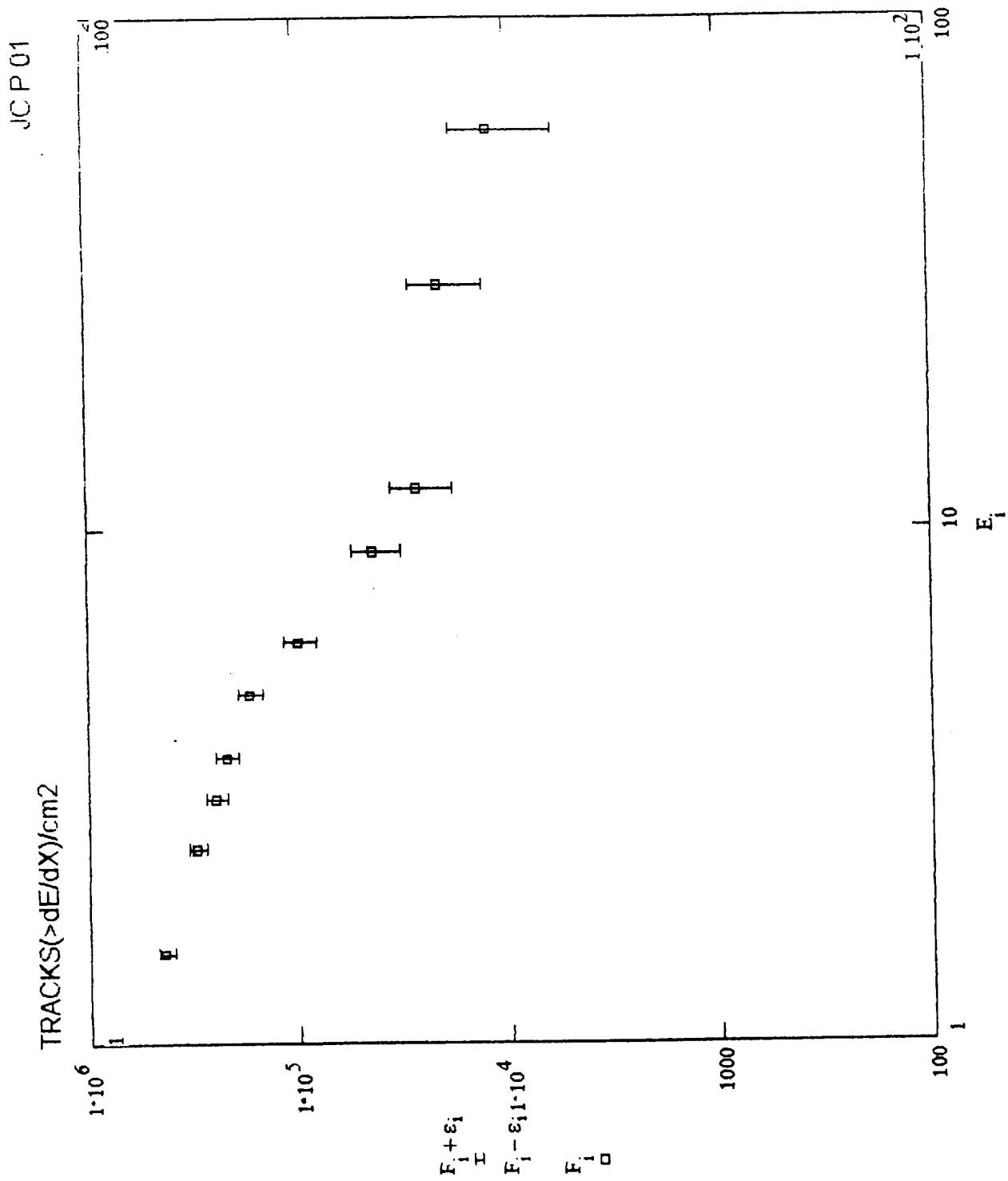
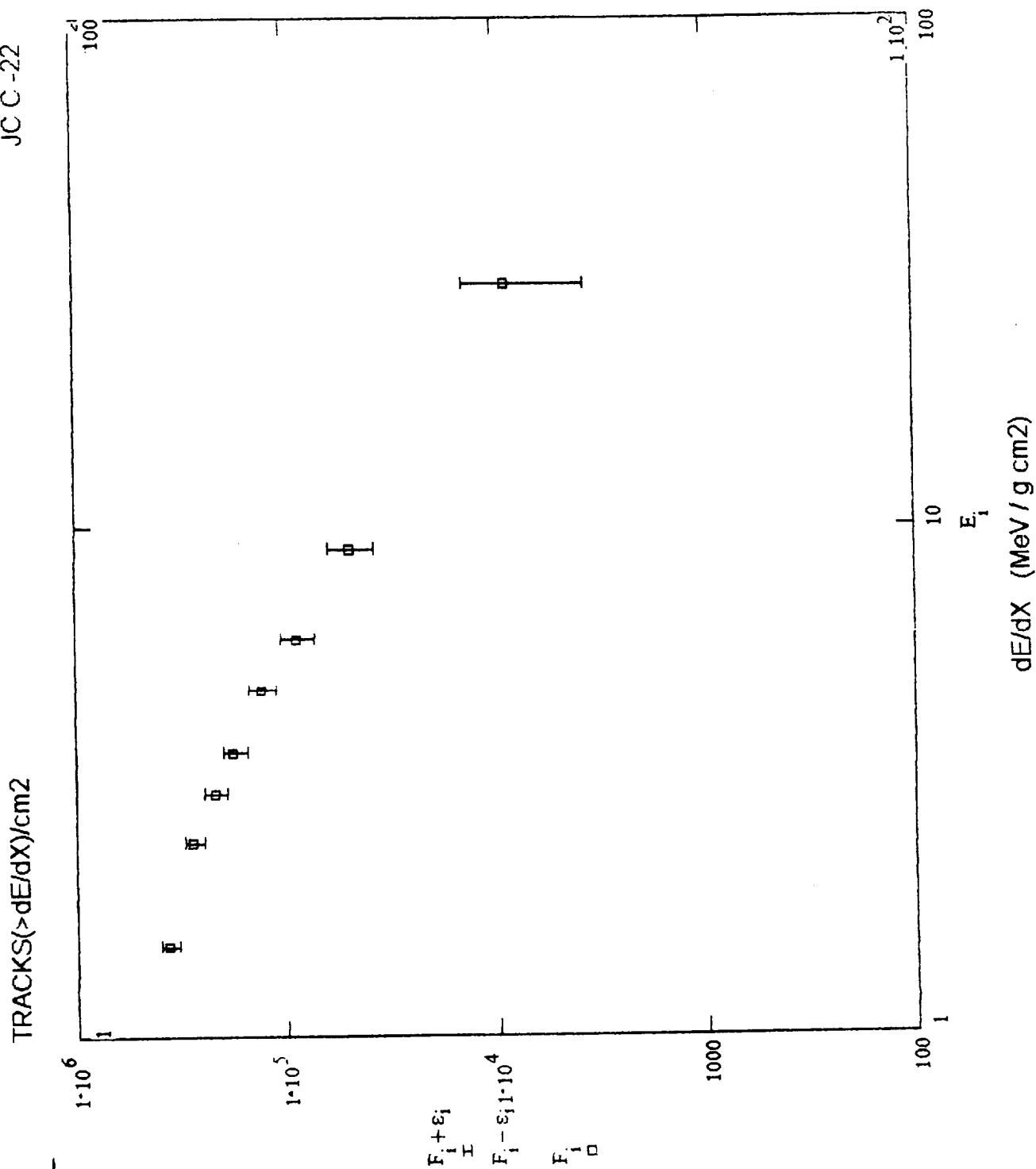


Figure 24-c Integral Linear Energy Transfer Spectrum
in the emulsion plate located 94.5 g/cm2
from the top of ECT chamber



dE/dX (MeV / g cm²)
 Integral Linear Energy Transfer Spectrum
 in the emulsion plate located about 1.5 g/cm²
 from the top of IACEE chamber

Figure 24d



Integral Linear Energy Transfer Spectrum
in the emulsion plate located about 70 g/cm2
from the top of JACEE chamber

Figure 24e

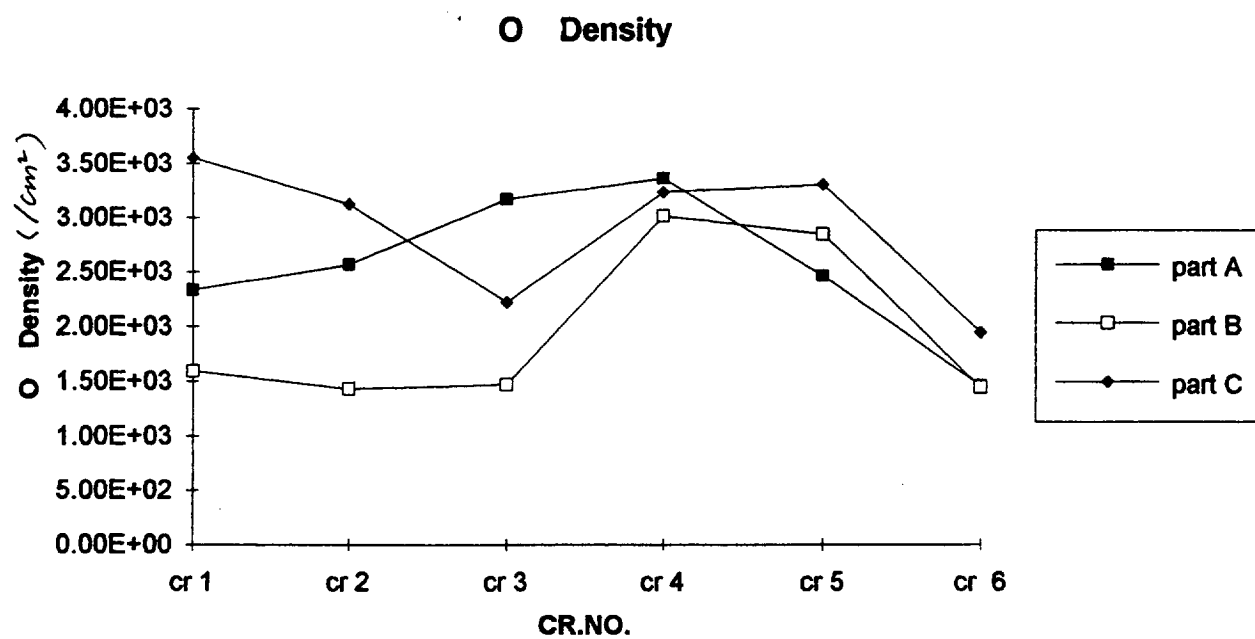


Figure 25a

O. Average Area

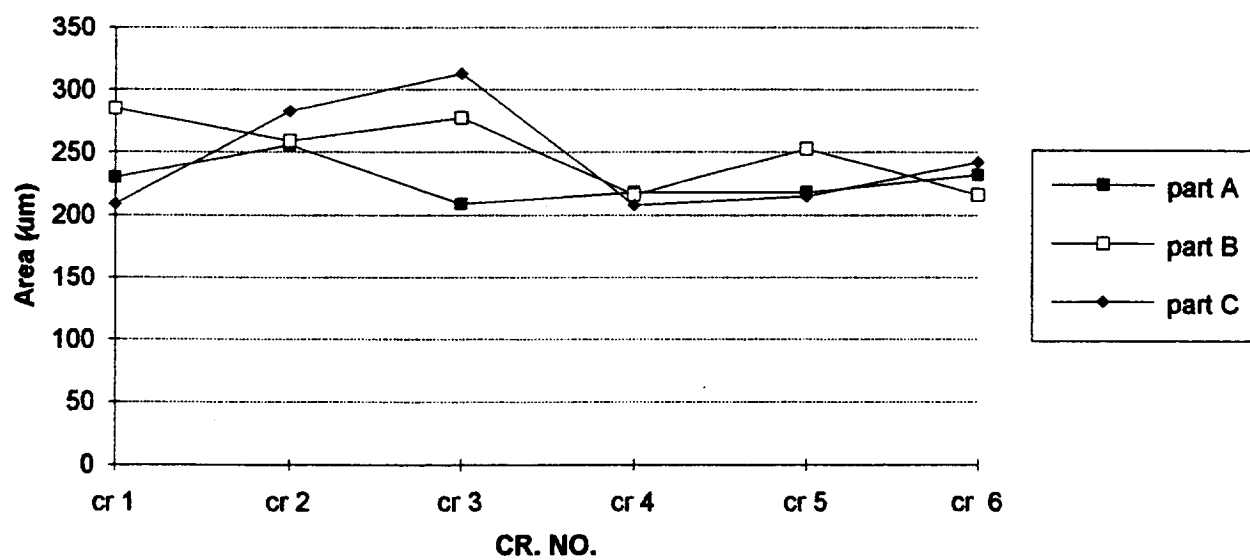


Figure 25b

CRJC12RR O.J.DS

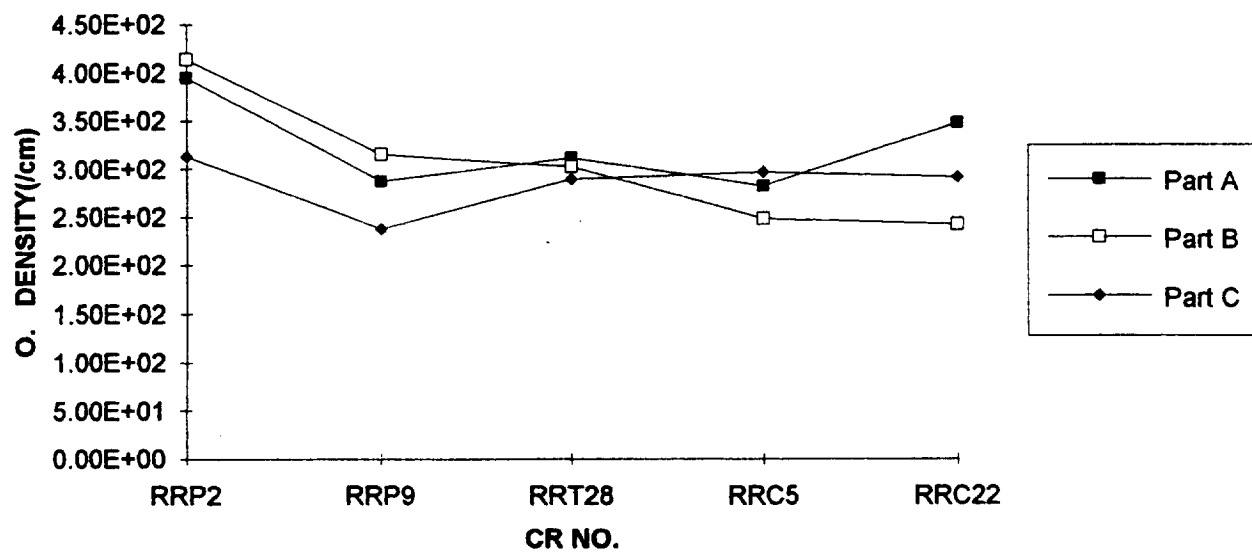


Figure 26a

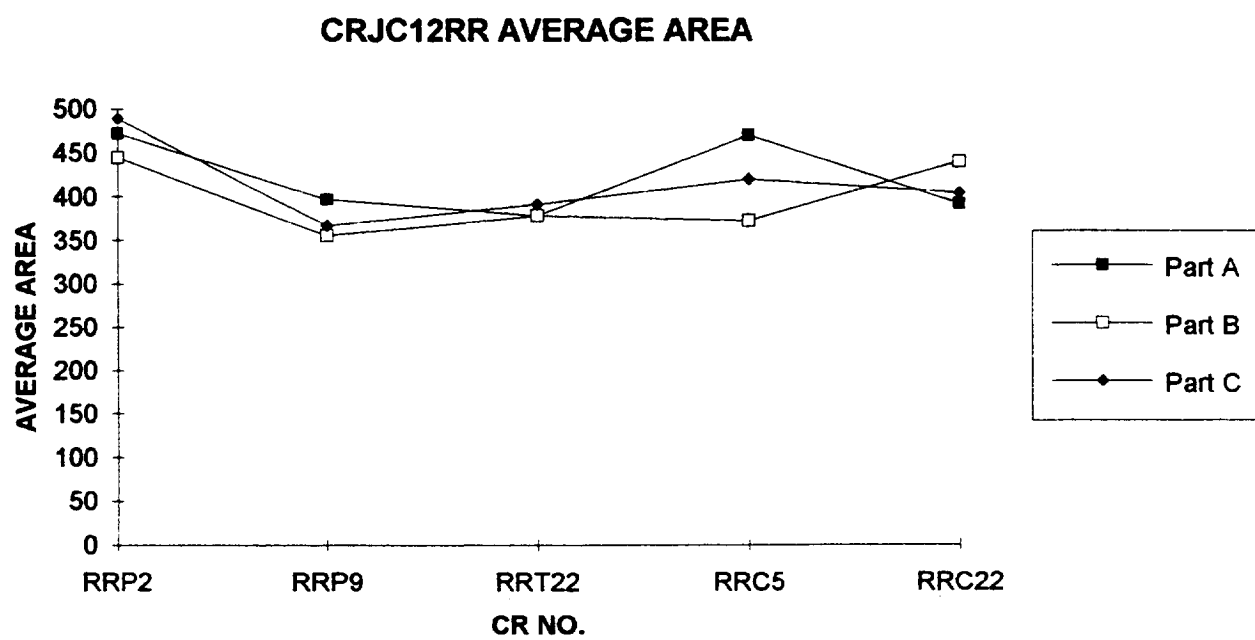


Figure 26b

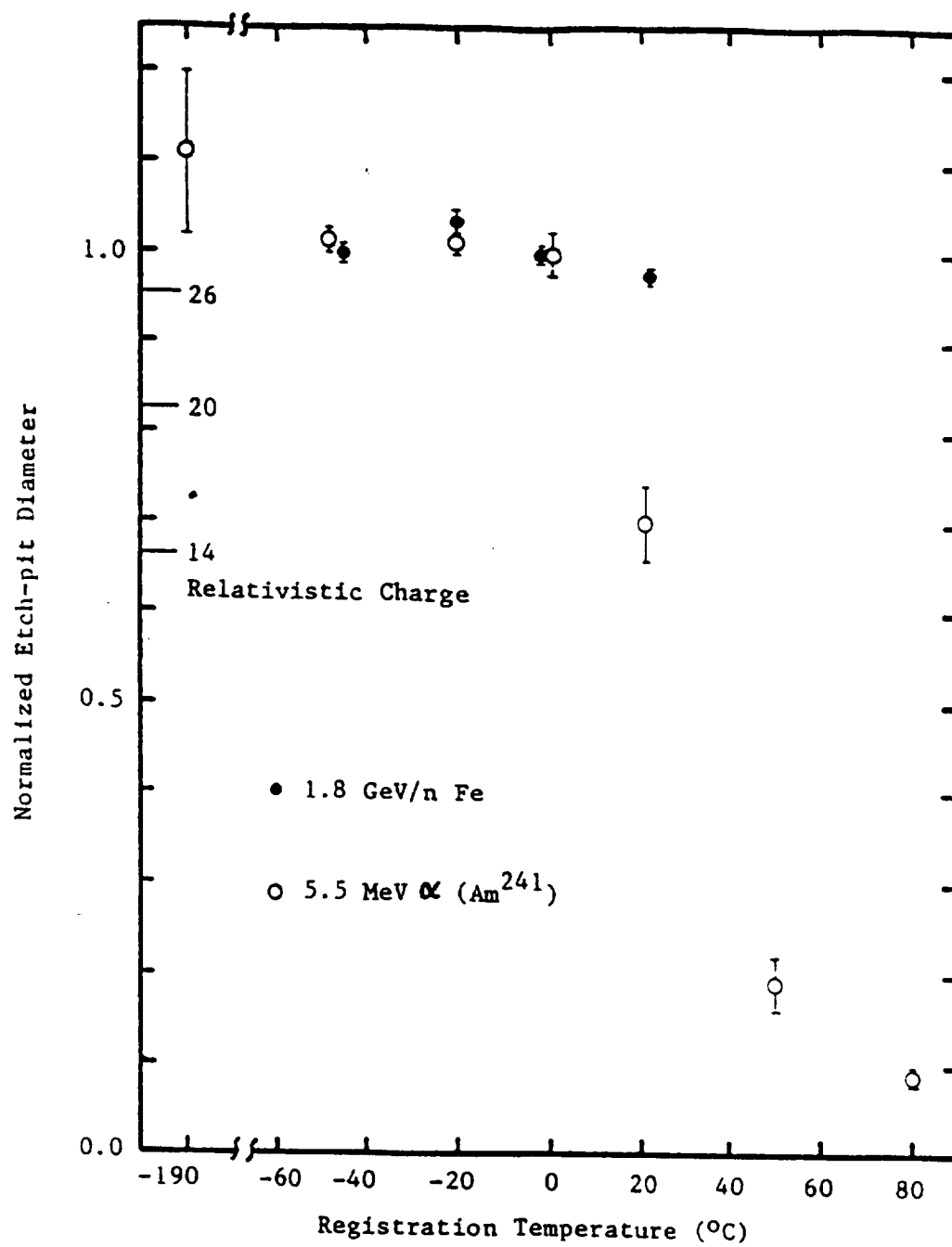


Figure 27

APPENDIX A

**ECT Dosimetry Report: LET Spectra,
Absorbed Doses and Neutron Dose Equivalents**

Measurement of LET Spectra, Absorbed Doses and
Low Energy Neutron Dose Equivalents in the
Emulsion Chamber Technology Experiment

Final Report

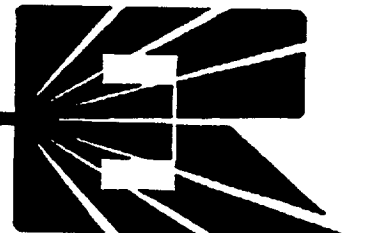
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28 August 1995

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Measurement of LET Spectra, Absorbed Doses and Low Energy Neutron Dose Equivalents in the Emulsion Chamber Technology Experiment

Abstract

LET spectra, total absorbed dose and low energy neutron fluence and dose equivalent measurements were made at various locations throughout the Emulsion Chamber Technology (ECT) experiment. LET spectra were measured under seven different shielding depths in the vertical center of the experiment stack. The low LET ($<10 \text{ keV}/\mu\text{m}$) region of the spectrum appeared to be dominated by stopping primary protons and was attenuated by the experiment shielding as measured from the lid to the base of the experiment stack. The mid ($>10 \text{ keV}/\mu\text{m}$) and high ($>100 \text{ keV}/\mu\text{m}$) LET regions of the spectrum appeared to be dominated by short range, high LET secondary particles produced in interactions between high energy protons and the nuclei of the experiment components. Only 20% of the high LET particles were long range ($>600 \mu\text{m}$) and thus considered to be galactic cosmic rays (GCR). Of this long range particles, $\sim 80\%$ were seen to be arriving from the direction of space (opposite Earth) and were stopping, illustrating the effect of the large amount of shielding in the ECT experiment on the GCR component. Little difference was seen in the overall integral LET flux spectra as a function of shielding.

Total absorbed dose was measured in TLDs at nine locations across the surface of each of the seven CR-39 PNTD pairs. Dose was seen to decrease as a function of shielding as measured from the top of the experiment stack. A mean dose of $238 \pm 3 \text{ mrad}$ was measured under $1.19 \text{ g}/\text{cm}^2$, while a mean dose of $142 \pm 1 \text{ mrad}$ was measured under $94.91 \text{ g}/\text{cm}^2$. There was little variation in dose amongst the different TLD locations across a given PNTD pair with the exception of PNTD Position 5. Dose through the experiment stack at Position 5 was substantially greater than that measured in other TLD positions, most likely due to the shielding environment surrounding the experiment and the orientation of the spacecraft relative to the South Atlantic Anomaly (SAA) during much of the mission.

Thermal ($<0.2 \text{ eV}$) and Resonance (0.2 eV – 1 MeV) neutron dose equivalents were measured at three locations within the experiment stack. A thermal dose equivalent of $0.30 \pm 0.0 \text{ mrem}$ and resonance dose equivalent of $11.3 \pm 5.6 \text{ mrem}$ were measured under the maximum shielding of $94.91 \text{ g}/\text{cm}^2$. Maximum thermal and resonance neutron dose equivalents of 0.79 ± 0.16 and $39 \pm 19 \text{ mrem}$, respectively, were measured in the middle of the stack under $41.84 \text{ g}/\text{cm}^2$. This was substantially higher than similar measurements made under lower shielding in previous STS missions, demonstrating the effect of shielding as a neutron moderator and thermalizer.

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1 Introduction

The Emulsion Chamber Technology (ECT) experiment consisted of a thick stack of radiation sensitive materials including nuclear emulsion, x-ray film, and CR-39 plastic nuclear track detector (PNTD). Interspersed within the experiment stack were seven doublets of CR-39 to measure the LET spectra above $\sim 5 \text{ keV}/\mu\text{m}$. Arrayed inside each CR-39 doublet were Thermoluminescent Detectors (TLDs) to measure total absorbed dose. Also included in three locations within the experiment were neutron detectors to measure thermal ($<0.2 \text{ eV}$) and resonance (0.2 eV – 1 MeV) neutron fluences and dose equivalents.

The ECT experiment flew in the cargo bay of the Space Shuttle on the STS-62 mission. STS-62 was launched on 4 March 1994 at 7:53 A.M. Central Standard Time. The mission lasted a total of 335.27 hours or 13.969 days. STS-62 had a mean altitude of 296 km and an orbital inclination of 39.0° . STS-62 was an Earth observation mission and the cargo bay was open toward the Earth, placing the bulk of the orbiter between the ECT experiment and space.

Exposure to ionizing radiation can be analyzed in terms of energy spectra or Linear Energy Transfer (LET) spectra. The LET of a particle is a measure of the change in the energy of the particle per unit path length and varies inversely with the energy of the particle. A particle's LET is believed to be of greater relevance than its energy in terms of significance to radiation sensitive materials and components and to radiobiology since LET is a measure of energy transferred to the surrounding medium through which the particle is traveling. CR-39 PNTDs measure the fluence (particle density per unit area, solid angle) and the LET of ionizing radiation of $\text{LET}_{\infty}\cdot\text{H}_2\text{O} \geq 5 \text{ keV}/\mu\text{m}$. Fluence and LET measurements can be combined to produce integral fluence and dose LET spectra. The fluence or dose from particles greater than a given LET on the y -axis is plotted as a function of LET on the x -axis. TLDs measure total absorbed dose directly and measurements are presented in units of mrad. Since TLDs are not capable of recording LET, it is not possible to convert the dose to dose equivalent.

The altitude and inclination of the spacecraft orbit affects the relative contribution of the different radiation components to the total dose. For low inclination orbits (28.5°) above

~ 300 km, the major source of dose will be from trapped protons in the South Atlantic Anomaly (SAA). A low altitude, higher inclination space shuttle mission, (~ 300 km) at 39.0° inclination, such as STS-62 will receive a lower dose from trapped protons in the SAA than will lower inclination (28.5°) missions. Exposure to trapped protons can be seen in LET spectra measurements as stopping primary protons in the low (<10 keV/ μ m) LET region and as short range, high LET secondary particles in the mid (>10 keV/ μ m) and high (>100 keV/ μ m) regions. Relative and absolute contributions from GCR are functions of orbital inclination with polar orbits receiving the greatest GCR contribution and equatorial orbits receiving a smaller GCR contribution. High inclination, lower altitude orbits for the space shuttle (57°) receive the major contribution to total dose from GCR. However, the orbital inclination of STS-62 (39.0°) is high enough to allow GCR to make a significant contribution to dose and LET spectra.

2 Experiment

2.1 Assembly of Experiment

The CR-39 layers to be included in the ECT experiment were cut to dimensions of 49.8×39.8 cm². Thirteen (13) detector squares of 3×3 cm² were cut into the larger sheets. These squares were used to measure LET spectra following the experiment. A portion of one edge of each of the squares was left uncut so that the square detector would remain attached to the larger sheet during the experiment. Twenty-seven (27) circles of 0.5 cm diameter were also cut into the detector sheets to accommodate TLDs. Figure 1 shows the location of the LET spectra squares and the TLDs. Each square was numbered as to location in the larger sheet. The three TLDs nearest each numbered square share that number. A unique identifying detector number, of the form ESS-NN, was scribed into each square in the lower right-hand corner opposite the uncut edge. Figure 2 shows the numbering convention for each PNTD square. E stands for ECT, SS denotes the number of the CR-39 sheet in the batch, and NN denotes the position of the PNTD square on the sheet. A number of the form ESS was scribed in the lower right hand corner of the large CR-39 sheet in order to identify it after the PNTD squares had been removed. The scribed side of each layer is the

Table 1: Composition and type of each CR-39 pair assembled for the ECT experiment.

Top Layer Number	Bottom Layer Number	TLD
E04	E01	✓
E12	E05	✓
E15	E14	✓
E19	E20	✓
E21	E24	✓
E33	E25	✓
E35	E39	✓
E16	E-C	
E30	E38	

top side.

The CR-39 sheets were assembled into pairs. The polyethylene protective layers were removed from each CR-39 surface. A layer of 8 μm kimfoil (polycarbonate) was placed between the two CR-39 layers of the pair as a separator. The layers were carefully aligned so that the PNTD squares would line up with each other in the two CR-39 sheets. The layers were secured together around the edges of the CR-39 sheets with 6 mil Teflon (3M) tape. Additional Teflon tape was placed around the perimeter of the TLD holes in order to keep the two sheets from buckling. A TLD-700 (^7LiF) chip was placed in each of the TLD holes. The TLD chips were covered with Whatman filter paper and taped into place with Teflon tape. The finished detector assembly was then covered in a protective layer of 25 μm thick polycarbonate foil. Nine (9) CR-39 pairs were assembled, seven (7) with TLDs and two (2) without TLDs. Table 1 lists the sheet number and assembly type of each of the CR-39 pairs.

Thermal and Resonance Neutron Detectors (TRNDs) consisting of layers of CR-39 PNTD and ^6LiF were assembled for inclusion in the ECT experiment. The TRNDs are roughly $1/2'' \times 1/2''$ and are ~ 3 mm in thickness. TRNDs were assembled in pairs with one being covered with a layer of Gd foil neutron absorber. The TRND pairs are covered on both sides by a layer of Whatman filter paper and are labeled. Three (3) flight and one (1) ground control TRND were assembled. Table 2 shows the contents and label of each TRND pair.

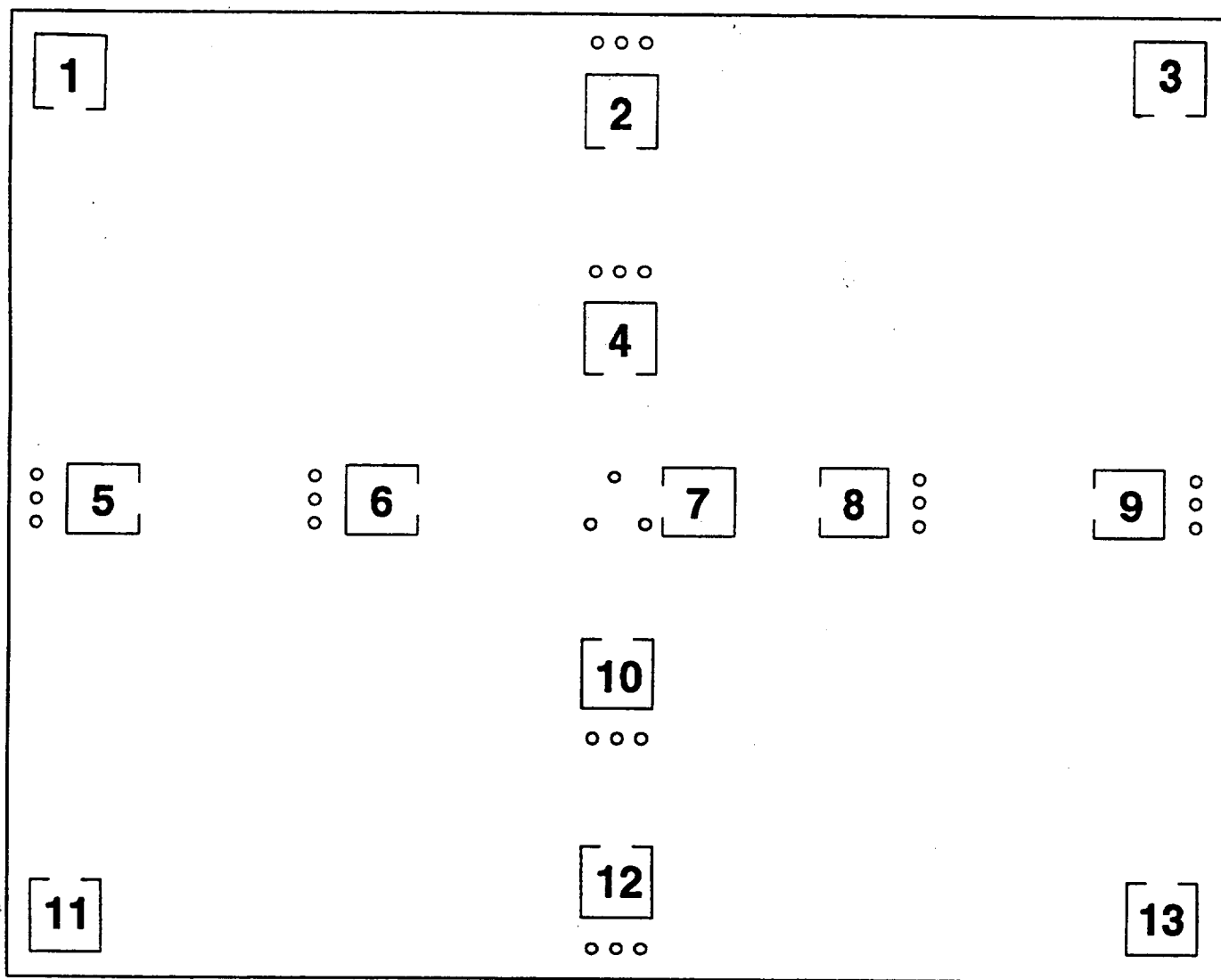


Figure 1: Positions of the CR-39 cut-outs and TLDs on the surface of the large CR-39 detector pairs.

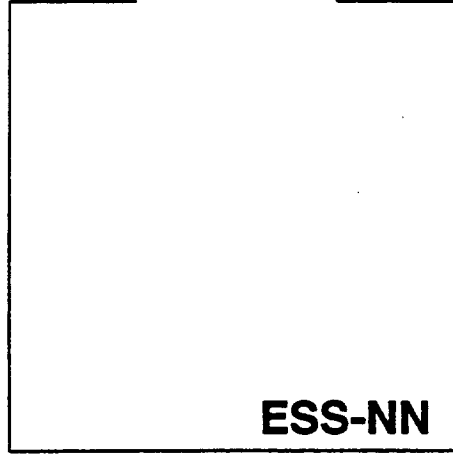


Figure 2: Numbering convention for CR-39 PNTD cut-outs. E stands for ECT, SS is the number of the CR-39 sheet, and NN is the position of the detector in the sheet.

Table 2: Thermal and Resonance Neutron Detectors (TRNDs) to be included in the ECT experiment.

Pair	Type	Gd foil
A1,A2	flight	A1
B1,B2	flight	B1
C1,C2	flight	C1
D1,D2	ground	D1

Table 3: Shielding of CR-39 and TLD layers relative to the normal of the detector stack. Shielding is measured relative to both the top and the bottom of the experiment.

Doublet No.	Top to Bottom Shielding (g/cm ²)	Bottom to Top Shielding (g/cm ²)
E33/E25	1.19	93.87
E21/E24	8.36	86.70
E15/E14	26.71	68.35
E19/E20	41.84	53.22
E34/E39	69.45	25.61
E12/E05	87.07	7.99
E04/E01	94.91	0.15

The assembled CR-39 pairs and TRNDs were then shipped to University of Alabama, Huntsville for integration into the ECT experiment. A set of 16 ground control TLDs of the same manufactured batch as those included in the CR-39 assemblies were included for control purposes along with four CR-39 pieces.

2.2 Shielding of the ECT Experiment

The ECT experiment consisted of a thick stack of radiation sensitive materials inside an aluminum canister and placed inside the Shuttle cargo bay. The orientation of the CR-39 sheets was normal to the orbiter vertical. Table 3 gives the shielding values of the CR-39 detectors and TLDs measured normal to the detector stack. During most of the mission, the orbiter was oriented upside-down relative to the Earth, that is the open cargo bay faced the Earth's surface. The shielding of the experiment can be measured from two origins, the top of the experiment closes to the Earth and the bottom of the experiment, closest to space. The first set of shielding values, top to bottom, are relevant in trying to understand the exposure of the experiment to low LET radiation including primary protons. The second set of shielding values, bottom to top, are of relevance in understanding the exposure of the experiment to GCR, since most of the GCR arrived in the experiment stack from the space side.

3 LET Spectra Measurements from CR-39 PNTDs

3.1 Calibration of CR-39 PNTDs

The CR-39 batch manufactured for the ECT experiment needed to be calibrated in order to transform measured track data from the experimental detectors into LET information. Samples of the ECT batch of CR-39 were exposed to heavy-ion beams of known energy and LET at the Lawrence Berkeley Laboratory Bevalac accelerator and to protons and α -particles at the University of Tokyo cyclotron. Samples of detector material were cut out of the ECT CR-39 sheets and machined into 4×4 cm² squares. These detectors were then exposed to a fluence of $\sim 2.5 \times 10^3$ particles/cm². The detectors were exposed in ambient pressure and temperature. Beam direction was normal to the detector surface.

Following exposure, the detectors were divided into two groups, one etched in a solution of 6.25 *N* NaOH at 50°C for 168 hours and the other etched in a solution of 6.25 *N* NaOH at 50°C for 36 hours. Bulk etch measurements were made by the loss-of-mass method. The detectors were read out using the Automated Track Analysis System. Track diameters were automatically measured in a 4 cm² region in the center of the detector. The reduced etch rate for each ion/energy pair was then calculated using the following formula:

$$V_R - 1 = \frac{1 + (d/2B)^2}{1 - (d/2B)^2} - 1, \quad (1)$$

where d is the mean track diameter and B is the bulk etch of the detector. A curve of the form:

$$x = a_0 + a_1y + a_2y^2 + a_3y^3 + a_4y^4, \quad (2)$$

was fitted to the calibration data points, where

$$x = \log_{10}(\text{LET}_{200} \cdot \text{CR} - 39) \quad (3)$$

and

$$y = \log_{10}(V_R - 1). \quad (4)$$

Separate calibration functions were determined for the 36 hour and 168 hour detectors. These calibration functions were later used to convert the measured track parameters from the flight detectors into values of LET.

3.2 Chemical Processing of CR-39

LET spectra was measured in CR-39 PNTDs at seven shielding depths within the ECT stack. Doublets were removed from the thirteen locations on the large CR-39 sheets. The upper sheet of each doublet was etched for 36 hours in a solution of 6.25 *N* NaOH at 50°C. The lower sheet in each doublet was etch for 168 hours in 6.25 *N* NaOH at 50°C. The two etching regimes were used in order to enhance tracks from two different components of the LET spectra. The values of bulk etch (thickness of material removed from each surface of the CR-39 detector by chemical processing) of detectors processed for 168 hours and 36 hours were $\sim 40 \mu\text{m}$ and $\sim 8 \mu\text{m}$, respectively.

3.3 Analysis of Processed Detectors

LET spectra was measured in each of the two layers from the doublet in position 7 on the large CR-39 sheet. A three pass, semi-automated track analysis system was used to scan the detector surface and measure the particle tracks. During the first pass, an etched CR-39 layer was placed on the microscope stage and all the objects on the detector surface (both tracks and noise) were automatically located. During the second pass, a human operator reviewed all the detected objects, separating particle tracks from spurious noise events. Finally, during the third pass, the operator measured semi-major and semi-minor axes of each elliptical track opening. The measured track parameters were converted to the unitless reduced etch rate ratio, V_R , by:

$$V_R = \sqrt{1 + \frac{4(a/B)^2}{[1 - (b/B)^2]^2}}, \quad (5)$$

where a and b are the semi-major and semi-minor axes of the track surface openings and B is the bulk etch of the detector. Values of V_R were converted to LET using the calibration functions determined for the specific manufactured batch of CR-39 employed in the ECT experiment. Separate calibration functions were used for the 168 and 36 hour etches.

3.4 Generation of LET Spectra

Two etching regimes were used in order to enhance tracks produced by particles belonging to different components of the LET spectra. The 168 hour etch allows low LET tracks to become sufficiently large to be easily located and measured. However, the long etch period over-etches tracks produced by short-range, high-LET secondaries. The 36 hour etch permits short-range, high-LET secondary particle tracks to be measured since these tracks are not over-etched during the short etch period. Low LET tracks are too small to be easily measured using such a short etch time.

The LET spectra for each layer of the doublet was determined separately. The spectra for the two etch durations were then combined to create a total LET spectrum. Figure 3 shows the integral LET flux spectra measured in the two detectors of the E04/E01 doublet for the 36 hour etch, the 168 hour etch and the combined spectra. At low LETs, there is little difference between the 168 hour spectrum and the combined spectra, while at high LETs, the 36 hour spectrum is identical to the combined spectra. The 168 hour spectrum falls off more rapidly than does the 36 hour spectrum at high LETs, illustrating that the longer etch period causes some of the high LET data to be lost. This data mostly takes the form of tracks from short-range, high-LET secondary particles. These particles have ranges less than the $\sim 40 \mu\text{m}$ bulk etch produced by the 168 hour etch, and the resulting tracks are over-etched and unmeasurable after this thickness of detector is removed. The 36 hour etch only extends down to $\sim 30 \text{ keV}/\mu\text{m}$, while the 168 hour spectra extends down to $\sim 5 \text{ keV}/\mu\text{m}$, the minimum LET registration threshold in CR-39. The shorter etching regime does not provide enough time for the low LET tracks to become large enough to be readily located and measured. By combining the LET spectra for the two etch regimes, an LET spectrum covering both LET regions is obtained.

The values of uncertainty in the LET spectra have been estimated. Figure 4 shows the combined integral LET flux spectrum for detectors E04/E01 with error bars. In the low LET region ($\text{LET}_{\infty} \cdot \text{H}_2\text{O} \approx 10 \text{ keV}/\mu\text{m}$) the uncertainty in the integral flux spectrum is $\sim 6\%$. For $\text{LET}_{\infty} \cdot \text{H}_2\text{O} \approx 100 \text{ keV}/\mu\text{m}$, the integral LET flux spectrum uncertainty is $\sim 8\%$. At high LET ($\text{LET}_{\infty} \cdot \text{H}_2\text{O} \approx 500 \text{ keV}/\mu\text{m}$), the uncertainty in integral flux spectrum is $\sim 40\%$. The

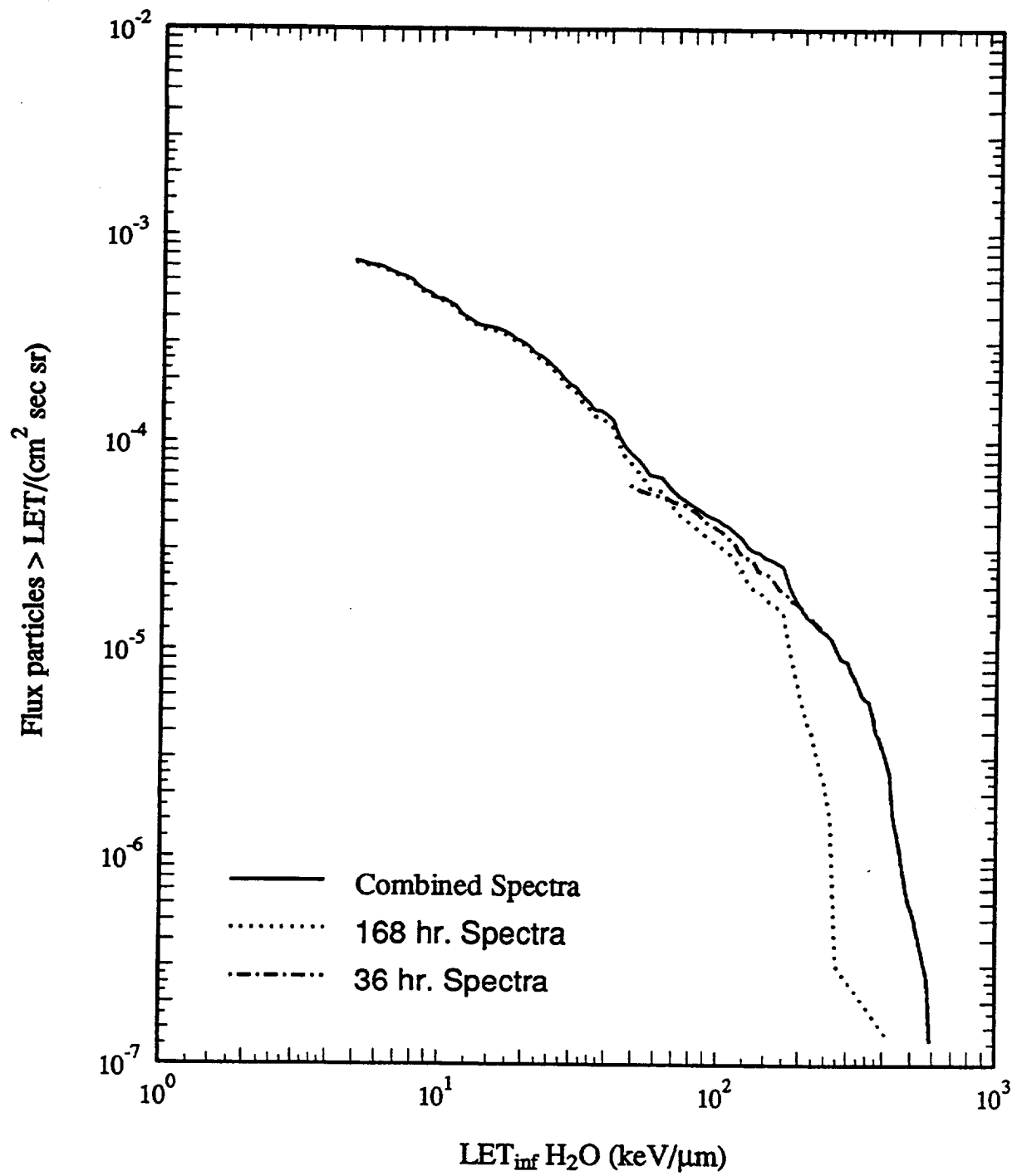


Figure 3: LET spectra measured in detector layer E01 etched for 36 hours, E04 etched for 168 hours, and E01 and E04 combined.

sizable increase in uncertainty at high LET is due to poor statistics since there are only a small number of tracks in this LET region. Values of uncertainty in integral LET flux spectra were similar for the other six measured LET spectra.

3.5 Results and Discussion of LET Spectra Measurements

Figure 5 shows the combined integral LET flux spectra for the seven CR-39 doublets measured for the ECT experiment. Figures 6 to 12 show separately the combined integral LET spectra for each doublet. Over most of the LET range, there is close agreement between all seven spectra within the limits of uncertainty. At low LET, the highest flux values were seen in detectors E33/E25 (Figure 6). This detector pair was closest to the lid of the experiment and was the least shielded pair from low LET particles arriving from the Earth direction. Since the cargo bay was opened toward the Earth and the layers E33/E25 were the most heavily shielded pair from particles arriving from the space (anti-Earth) direction, this detector pair had one of the lowest flux spectra in the high LET region ($\text{LET}_{\infty} \cdot \text{H}_2\text{O} > 100 \text{ keV}/\mu\text{m}$). A significant fraction of the particles in the high LET region are GCR and the largest flux of GCR is expected from the space direction.

Detector pair E04/E01 (Figure 12) was at the bottom of the ECT experiment stack, closest to the floor of the cargo bay. It was the most heavily shielded detector pair from particles arriving from the Earth direction and the least shielded from particles arriving from the space direction. E04/E01 had amongst the lowest flux values in both the low and high LET regions showing the complexity of shielding effects. In addition to the shielding from the experiment stack, there is also significant shielding from the orbiter itself, especially from the floor of the cargo bay which was parallel to the experiment stack.

The highest values for LET flux spectra in the mid and high LET regions ($\text{LET}_{\infty} \cdot \text{H}_2\text{O} \gtrsim 20 \text{ keV}/\mu\text{m}$) were measured in detector pairs E15/E14 and E19/E20 (Figures 8 and 9, respectively). These detector pairs were in the middle of the experiment stack. Given the shielding of these detector pairs from the experiment stack as measured from the space-end of the experiment, plus the shielding of the orbiter, these two detector pairs were probably the most heavily shielded of the experiment. This relatively large amount of shielding probably led

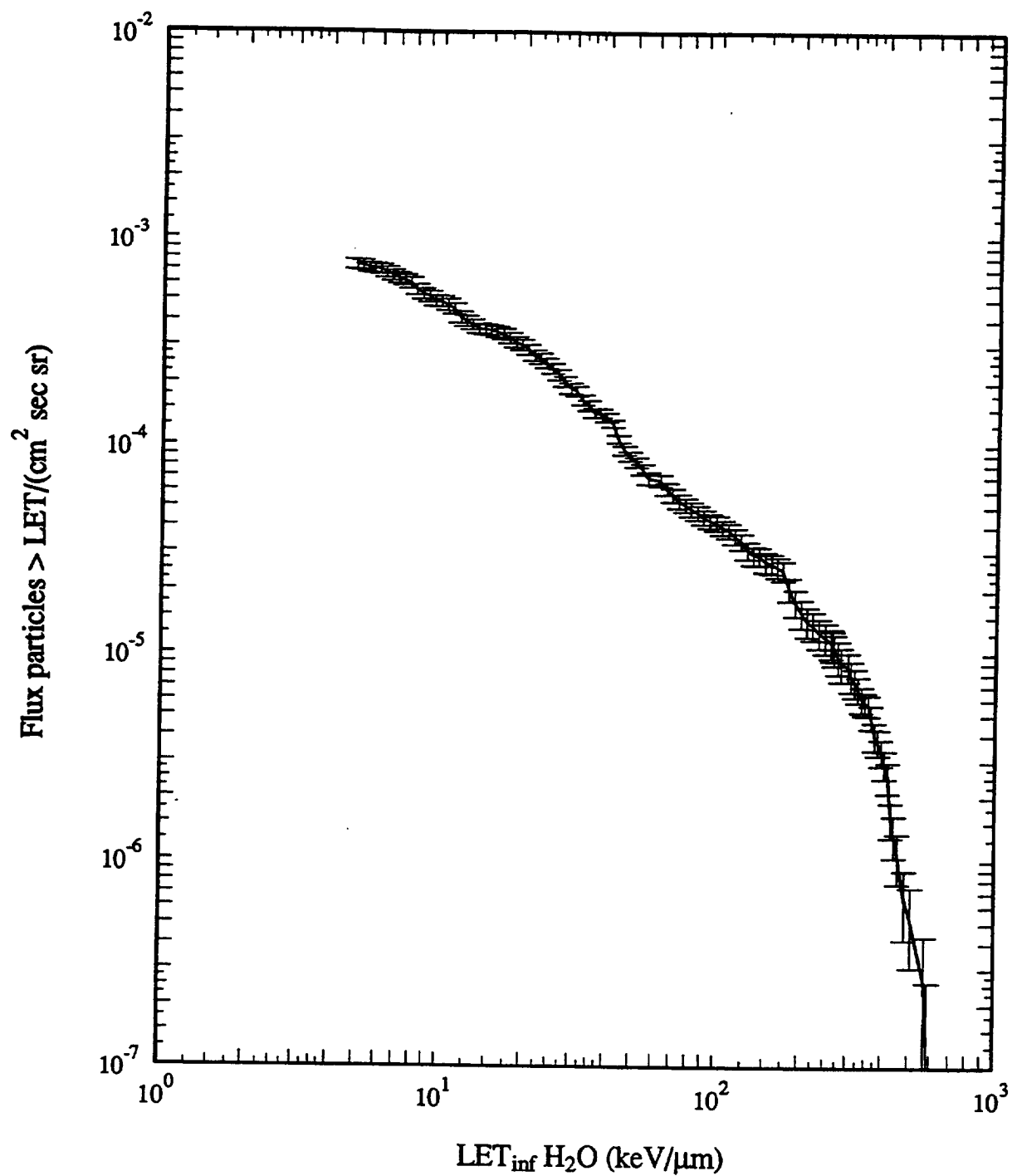


Figure 4: Combined integral LET flux spectra from detectors E04 and E01 with error bars illustrating uncertainty in the measurement. Uncertainty in the other six measured LET spectra was similar.

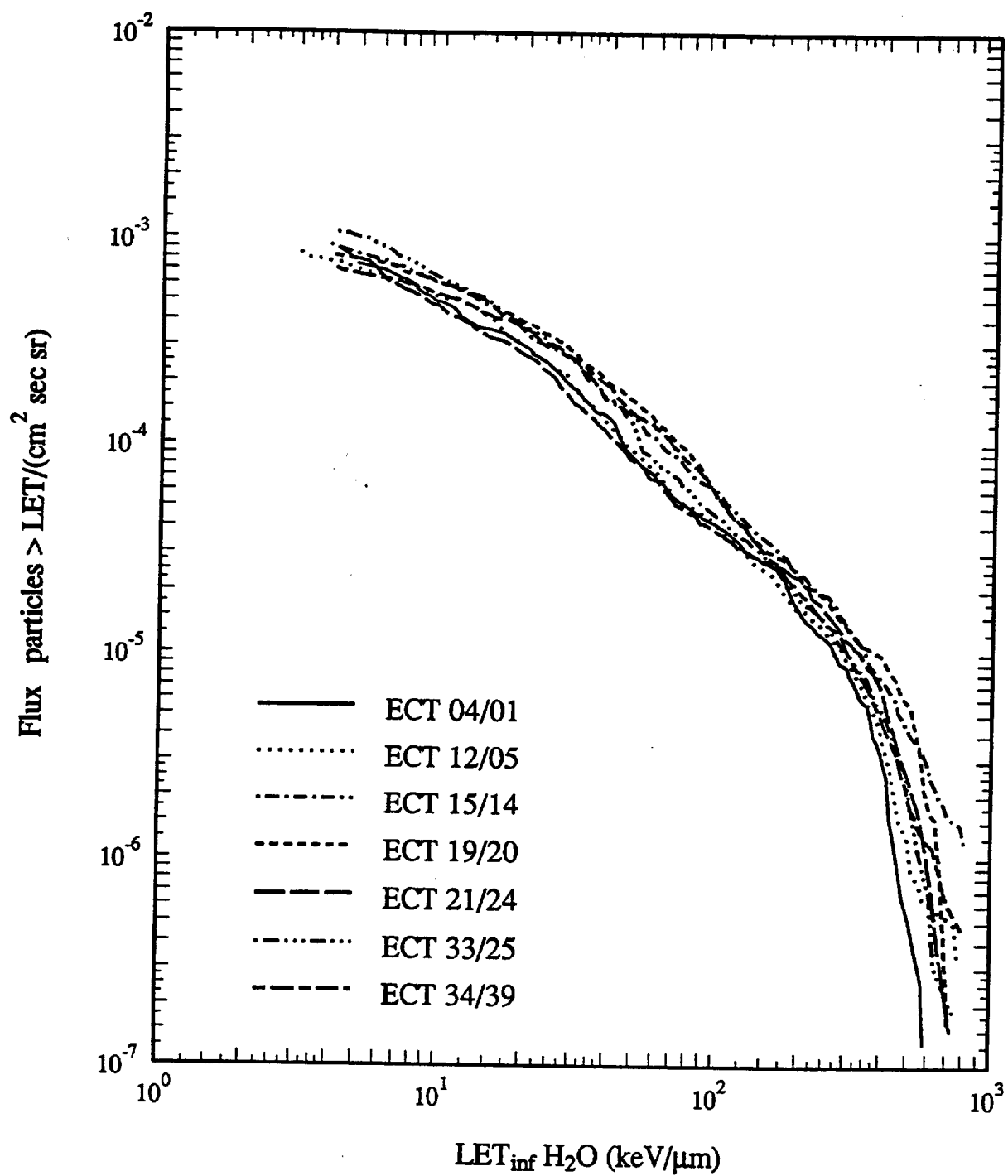


Figure 5: Combined Integral LET flux spectra for the seven CR-39 detector pairs measured in the ECT experiment.

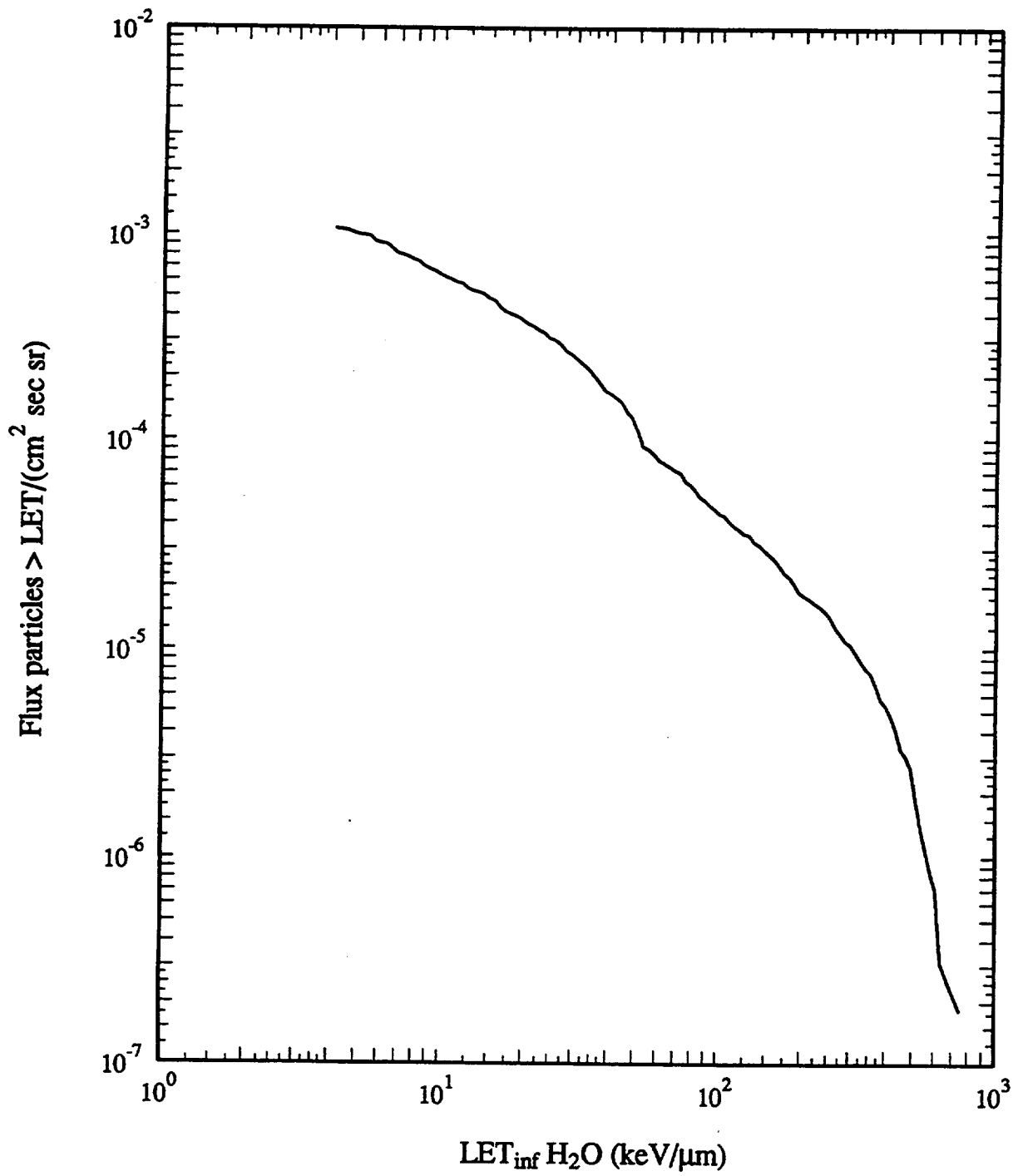


Figure 6: Combined Integral LET flux spectra for CR-39 detector pair E33/E25, under 1.19 g/cm² shielding, measured from the lid of the experiment container.

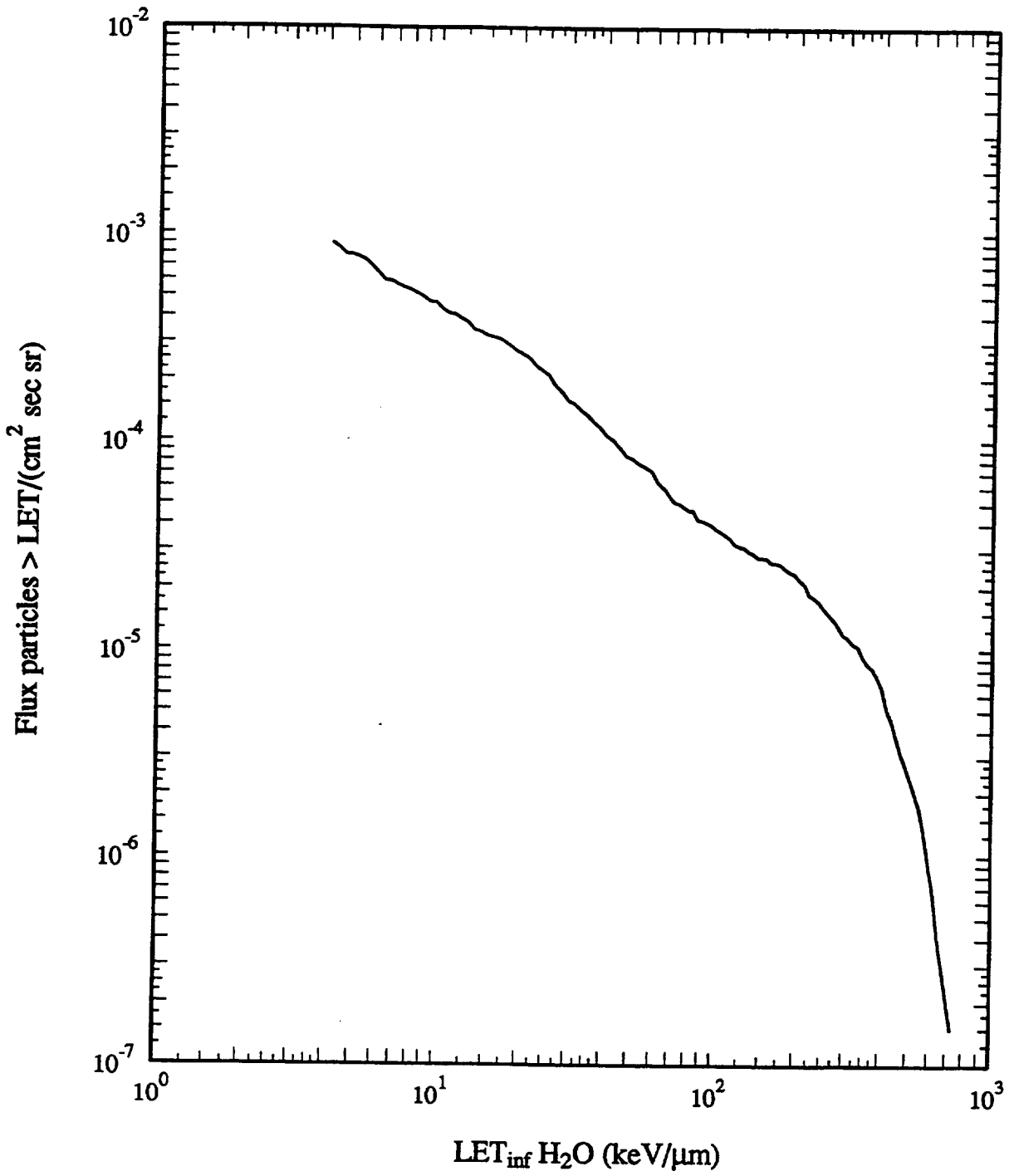


Figure 7: Combined Integral LET flux spectra for CR-39 detector pair E21/E24, under $8.36 \text{ g}/\text{cm}^2$ shielding, measured from the lid of the experiment container.

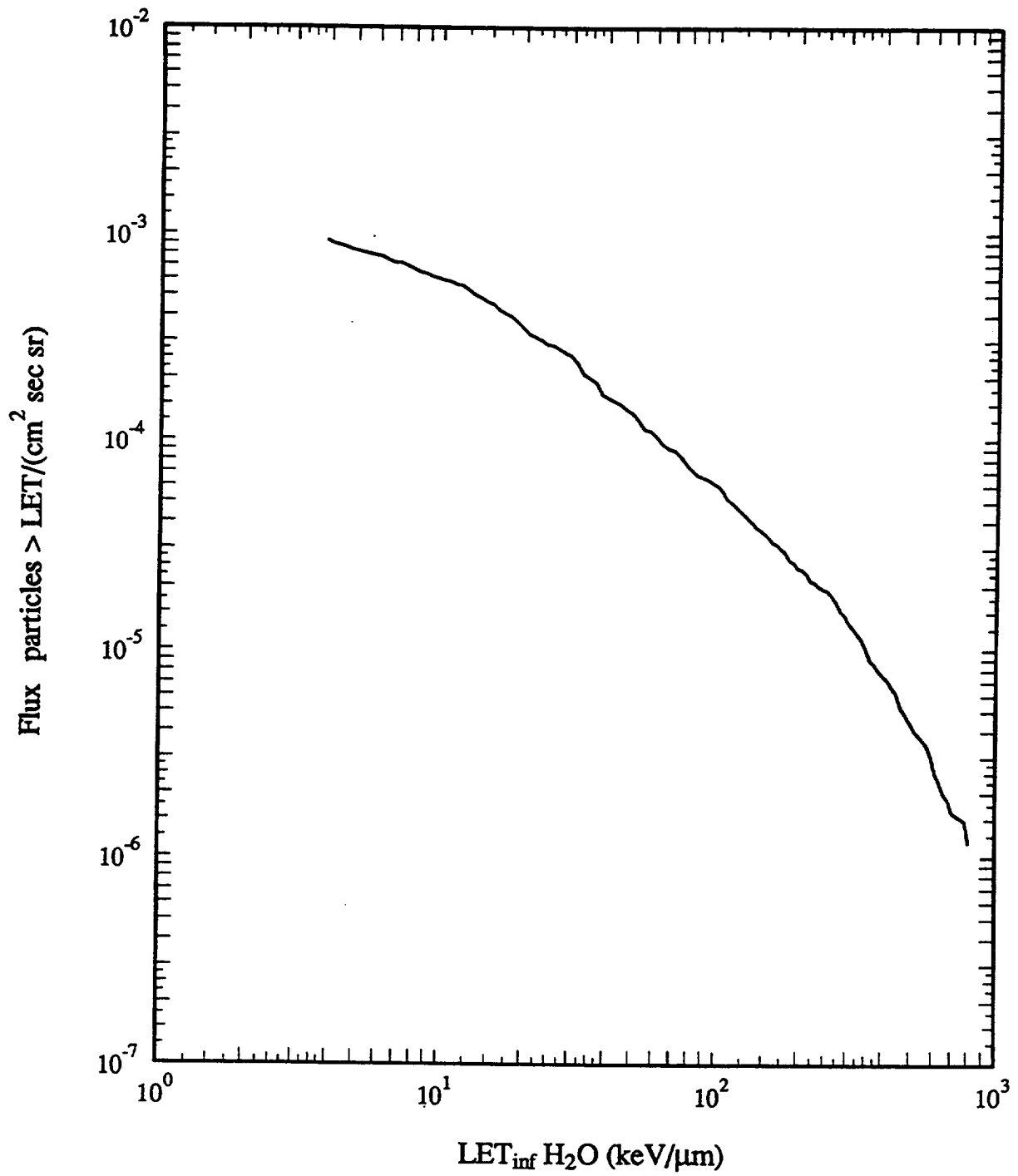


Figure 8: Combined Integral LET flux spectra for CR-39 detector pair E15/E14, under $26.71 \text{ g}/\text{cm}^2$ shielding, measured from the lid of the experiment container.

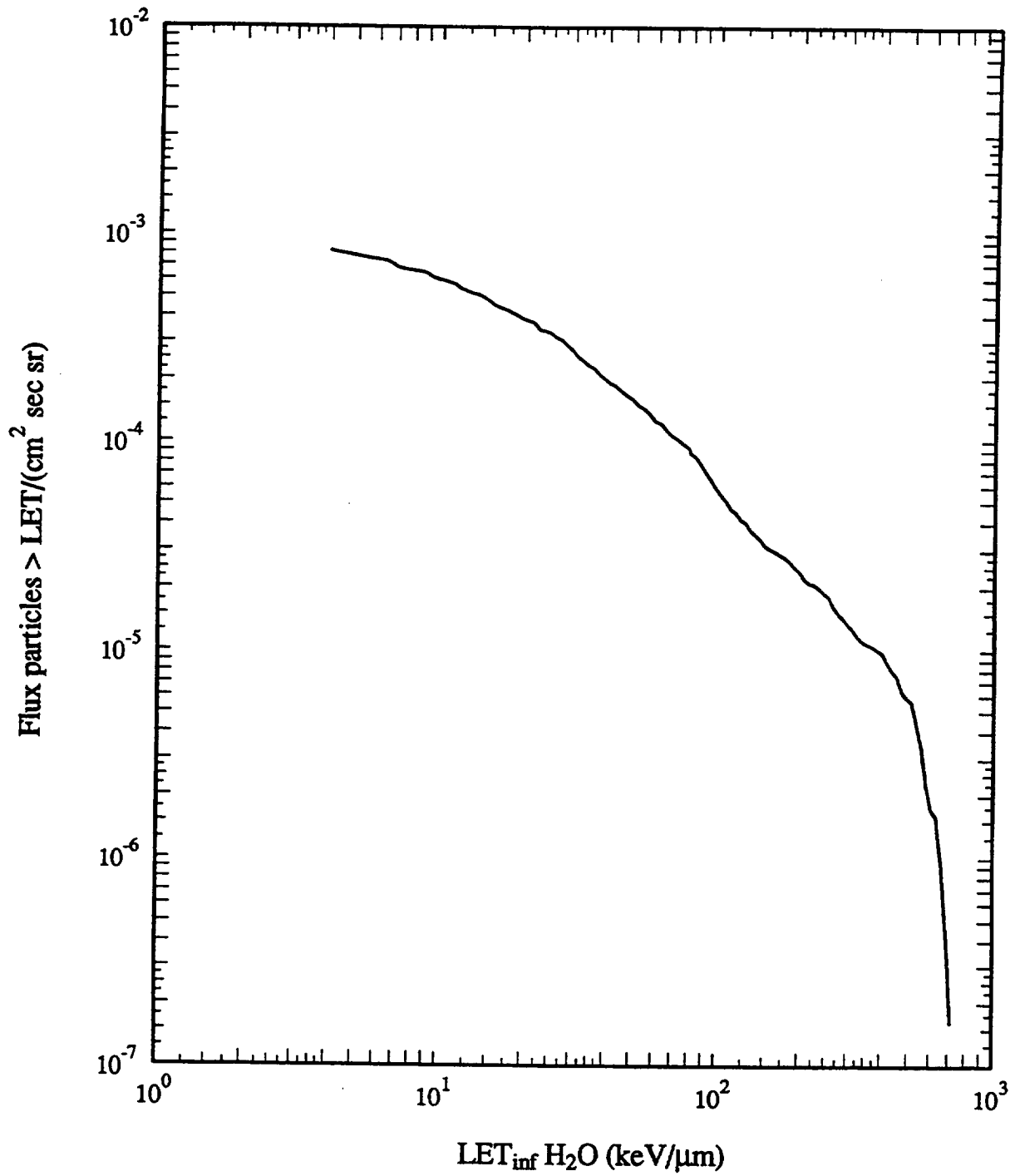


Figure 9: Combined Integral LET flux spectra for CR-39 detector pair E19/E20, under 41.84 g/cm² shielding, measured from the lid of the experiment container.

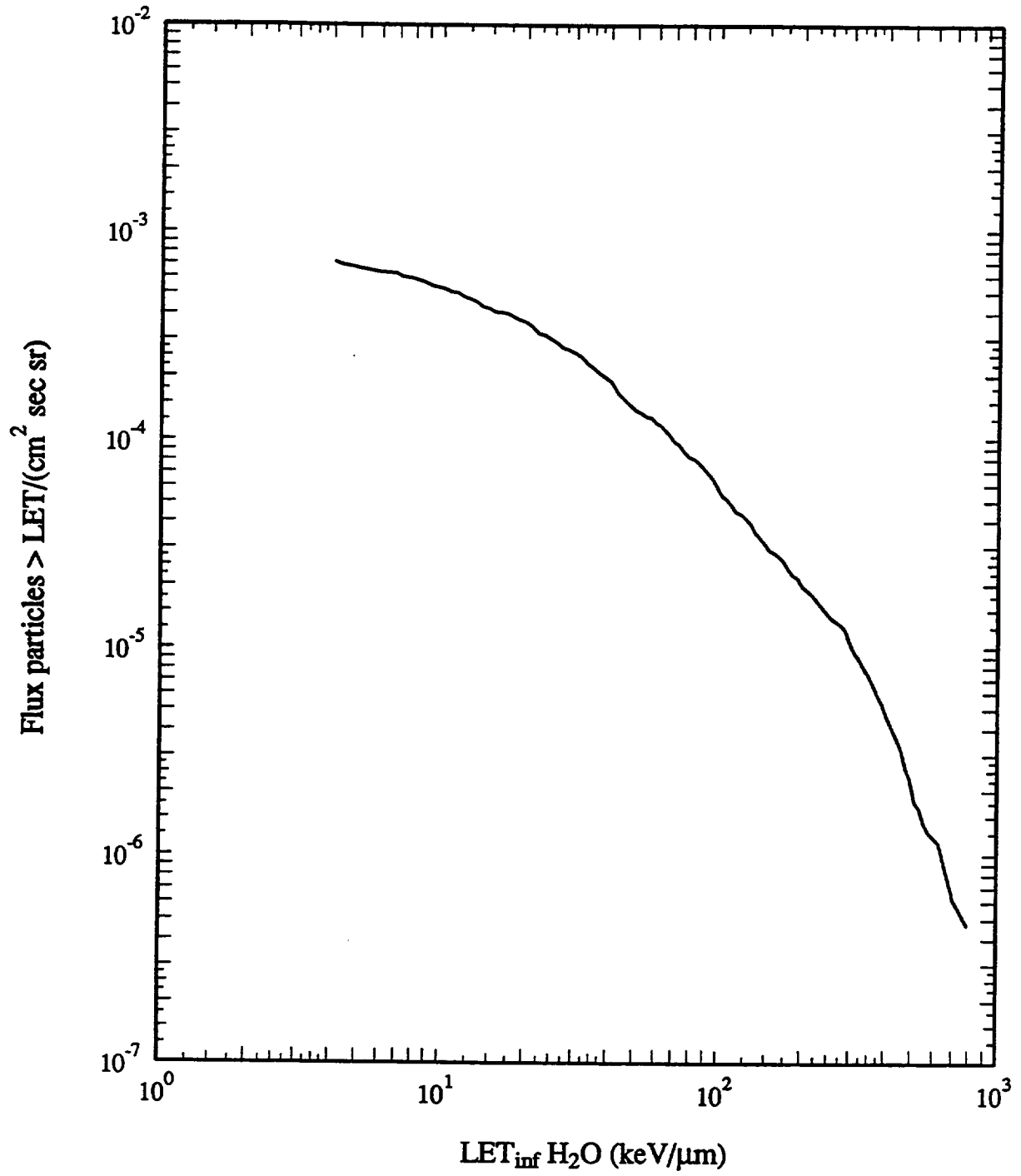


Figure 10: Combined Integral LET flux spectra for CR-39 detector pair E34/E39, under 69.45 g/cm^2 shielding, measured from the lid of the experiment container.

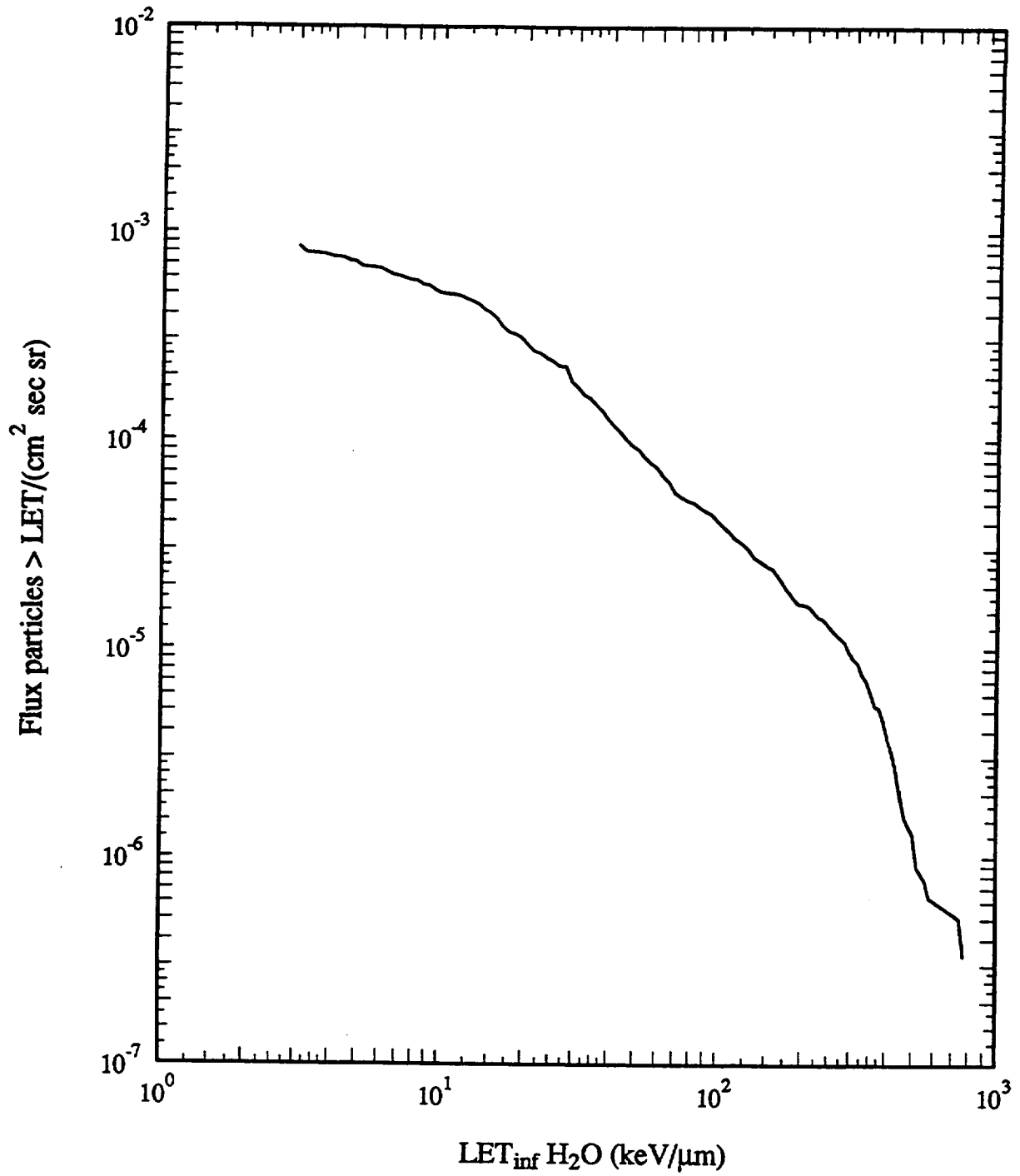


Figure 11: Combined Integral LET flux spectra for CR-39 detector pair E12/E04, under 87.07 g/cm² shielding, measured from the lid of the experiment container.

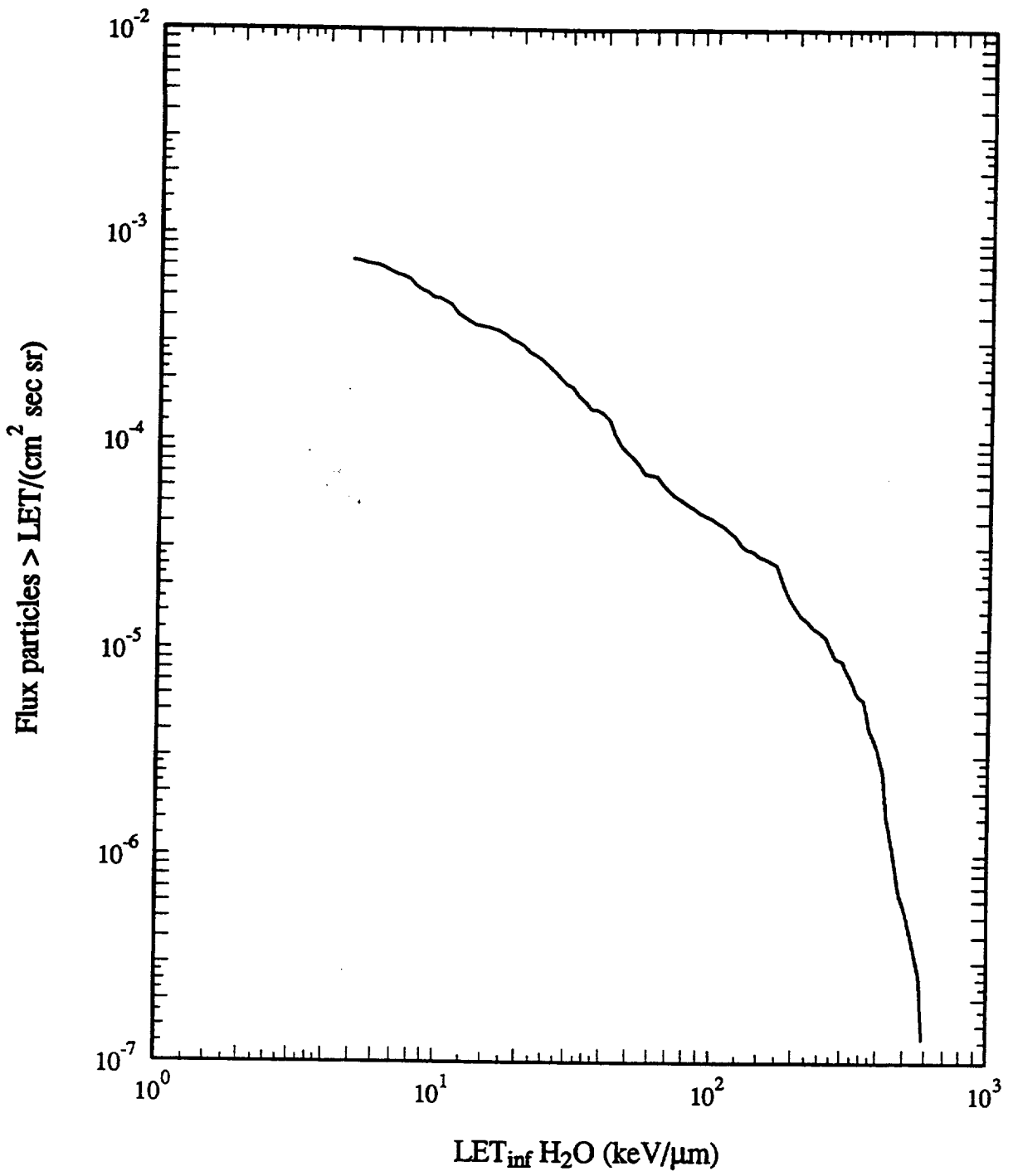


Figure 12: Combined Integral LET flux spectra for CR-39 detector pair E04/E01, under 94.91 g/cm² shielding, measured from the lid of the experiment container.

to a large production of short-range, high-LET secondary particles in these layers. Particles passing through the experiment were reduced in energy and increased in cross sections from nucleon interactions. They subsequently underwent inelastic collisions with the heavier nuclei (C and O) of the CR-39 and surrounding materials, producing the relatively high signal measured in the mid to high LET region.

Particle events were divided into two categories, long range events ($>600\ \mu\text{m}$) consisting mostly of GCR, and short range events ($<600\ \mu\text{m}$). Long range particle events were identified when two tracks, one of each surface of the detector layer, were found to have been made by the same particle. Since the pre-etch thickness of the CR-39 layers was $\sim 600\ \mu\text{m}$, the minimum range of two surface particles events was on the order of this thickness. The remaining particle events took the form of single tracks on only one surface of the CR-39 detector. At low LETs, ($<10\ \text{keV}/\mu\text{m}$), most of these events are made up of stopping primary protons. At higher LETs, a significant fraction of these events are caused by short-range, high-LET secondary particles. The minimum range of a particle observed in CR-39 as a track following a 36 hour etch is $\sim 8\ \mu\text{m}$.

For all seven detector pairs measured, $\sim 20\%$ of the particle events were long range. In CR-39 detectors flown in less heavily shielded configurations (i.e. inside a Shuttle mid-deck locker), long range events show little change in LET through the thickness of the detector. Tracks on each of the two detector surfaces are roughly of equal size, indicating that the particle's LET is changing only slowly. For a majority of the long range particle events measured in the ECT CR-39 layers, the track on the surface of the detector facing space was often smaller than the track on the surface facing Earth. This was seen in $\sim 80\%$ of the long range particle events and indicates that the particles were arriving from the direction of space and that they were near the end of their ranges (i.e. the change in track size indicates the rapid change in LET seen near the particle's Bragg peak). That fact that so many of the long range particle events were near their stopping points illustrates the effect of heavy shielding of the ECT experiment on the GCR component.

4 Dose Measurements with TLDs

Sets of three TLDs were positioned in nine locations across the plane of each of the seven CR-39 detector pairs (see Figure 1). Following disassembly of the detectors after flight, all the TLDs were read out along with background detectors. Measured dose values from each of the three TLDs within a given set were averaged together to obtain a mean dose for that particular position on a given CR-39 detector layer. Mean measured doses from TLDs are listed in Table 4, along with the uncertainty in each measurement, in terms of CR-39 pair and TLD position. Also listed in Table 4 is the mean dose and its uncertainty for all TLD positions on each CR-39 doublet. Figure 13 shows the measured TLD doses for each of the nine positions as a function of shielding depth measured from the bottom to the top of the experiment stack. Figure 14 shows a similar plot of dose as a function of shielding measured from the top of the stack to the bottom. Figure 15 shows the mean dose for each detector layer as a function of shielding depth measured from top to bottom.

As can be seen in Figures 14 and 15, dose dropped rapidly as a function of shielding between the first two layers, E33/E25 and E21/E24, from 248 ± 3 to 197 ± 2 mrad within ~ 7 g/cm². Dose remained relatively constant between layers E21/E24 and E12/E05, dropping from 197 ± 2 to 181 ± 2 mrad through ~ 79 g/cm². The dose again fell off rapidly between layers E12/E05 and E04/E01, dropping from 181 ± 2 to 142 ± 1 mrad in ~ 8 g/cm².

Measured doses in each of the nine TLD positions were relatively similar across a given detector layer. Only TLD Position 5 differed significantly from this pattern. The initial decrease in dose between layers E33/E25 and E21/E24 is not as great as that measured in the other TLD locations and dose can be seen to rise between layers E21/E24 and E19/E20 at Position 5, while it remains relatively constant for the other locations. Between E19/E20 and E04/E01, dose at Position 5 falls off gently. The reason for this difference in dose as a function of shielding between TLD Position 5 and the other eight TLD locations is most likely due to a combination in differences in shielding and in the orientation of the spacecraft during most of the mission. TLD Position 5 was on the right-most side of the detector sheet as can be seen in Figure 1. Thus it was close to the vertical side of the experiment

Table 4: Tissue absorbed doses and uncertainties (mrad) for the nine TLD positions on each layer and for the seven CR-39 PNTD layers. Also included are average dose values for each layer.

CR-39 Doublet Number	Total Absorbed Dose (mrad)											
	TLD Position											
	2	4	5	6	7	8	9	10	12	Mean		
E33/E25	248 ± 7	250 ± 8	235 ± 7	258 ± 8	251 ± 8	275 ± 8	235 ± 7	266 ± 8	212 ± 6	248 ± 3		
E21/E24	195 ± 6	199 ± 6	223 ± 7	191 ± 6	191 ± 6	203 ± 6	188 ± 6	194 ± 6	188 ± 6	197 ± 2		
E14/E14	203 ± 6	198 ± 6	237 ± 7	198 ± 6	197 ± 6	197 ± 6	184 ± 6	176 ± 5	186 ± 6	197 ± 2		
E19/E20	177 ± 5	175 ± 5	242 ± 7	180 ± 5	175 ± 5	167 ± 5	176 ± 5	171 ± 5	182 ± 5	193 ± 2		
E34/E39	185 ± 6	183 ± 5	230 ± 7	186 ± 6	171 ± 5	166 ± 5	179 ± 5	176 ± 5	195 ± 6	186 ± 2		
E12/E05	177 ± 5	174 ± 5	207 ± 6	177 ± 5	169 ± 5	180 ± 5	173 ± 5	175 ± 5	195 ± 6	181 ± 2		
E04/E01	143 ± 4	144 ± 4	175 ± 5	138 ± 4	135 ± 4	125 ± 4	124 ± 4	136 ± 4	154 ± 5	142 ± 1		

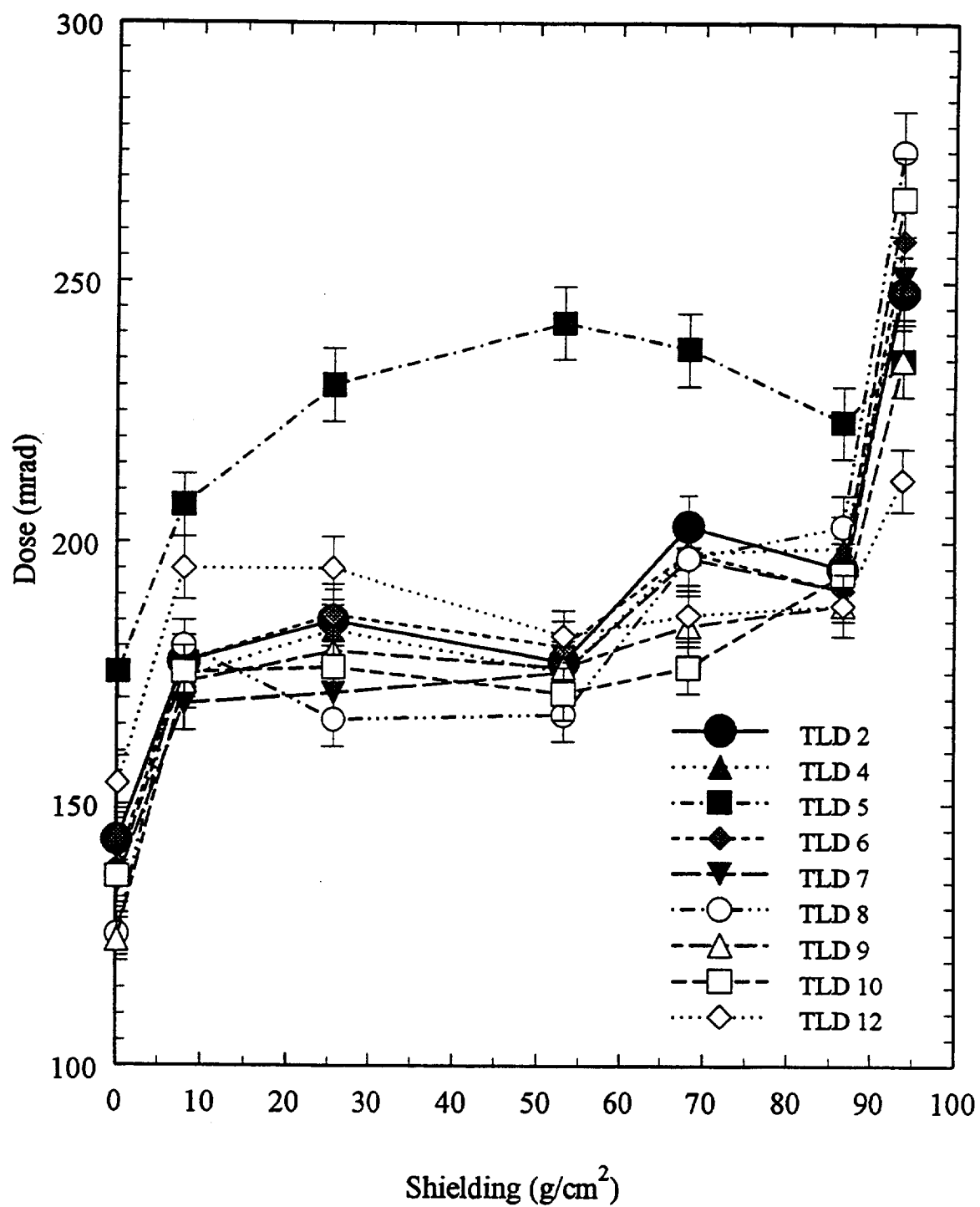


Figure 13: Dose in TLDs as a function of shielding depth as measured from the bottom to the top of the experiment stack for each of the nine TLD positions.

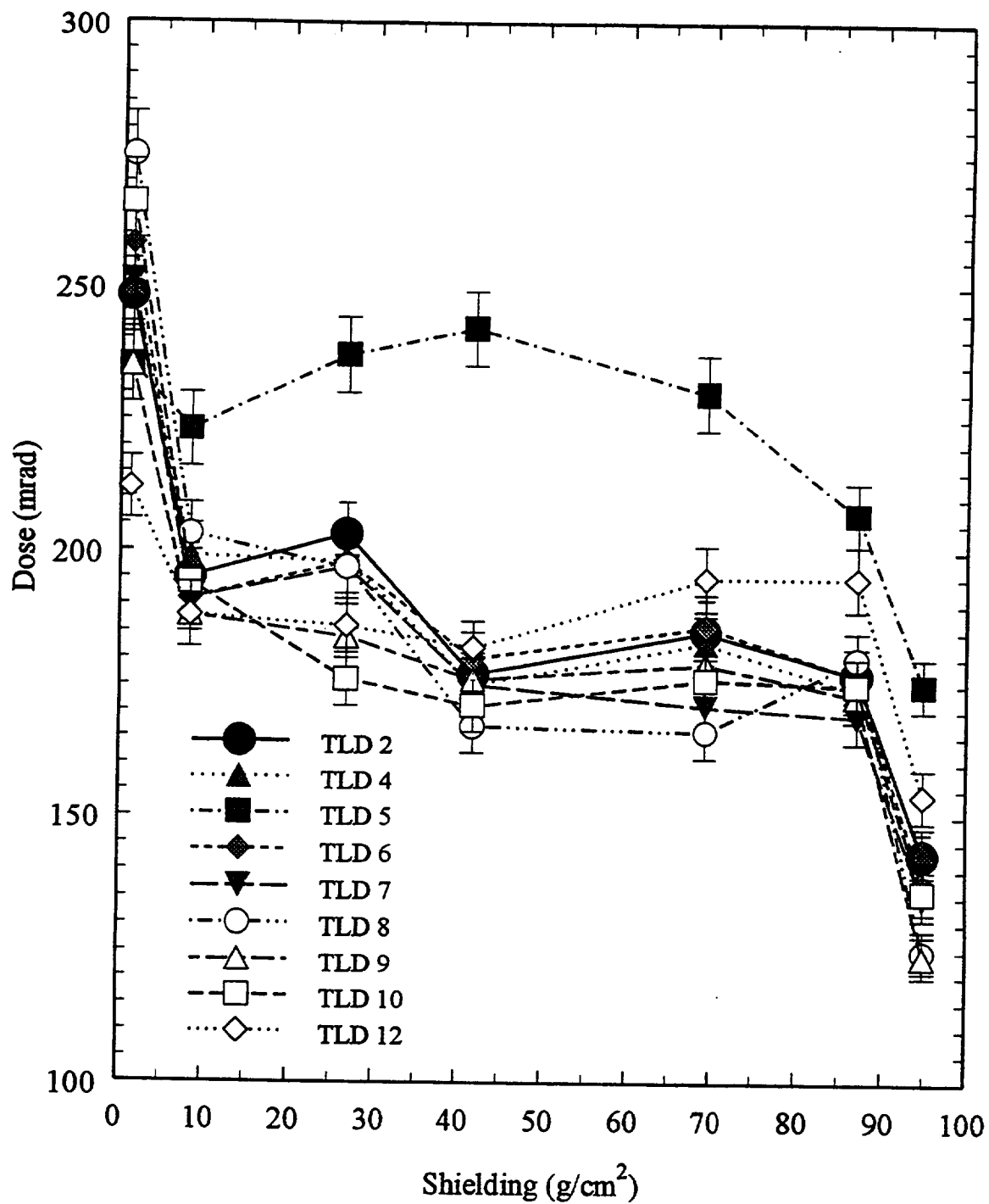


Figure 14: Dose in TLDs as a function of shielding depth in the experiment stack as measured from top to bottom for each of the nine TLD positions.

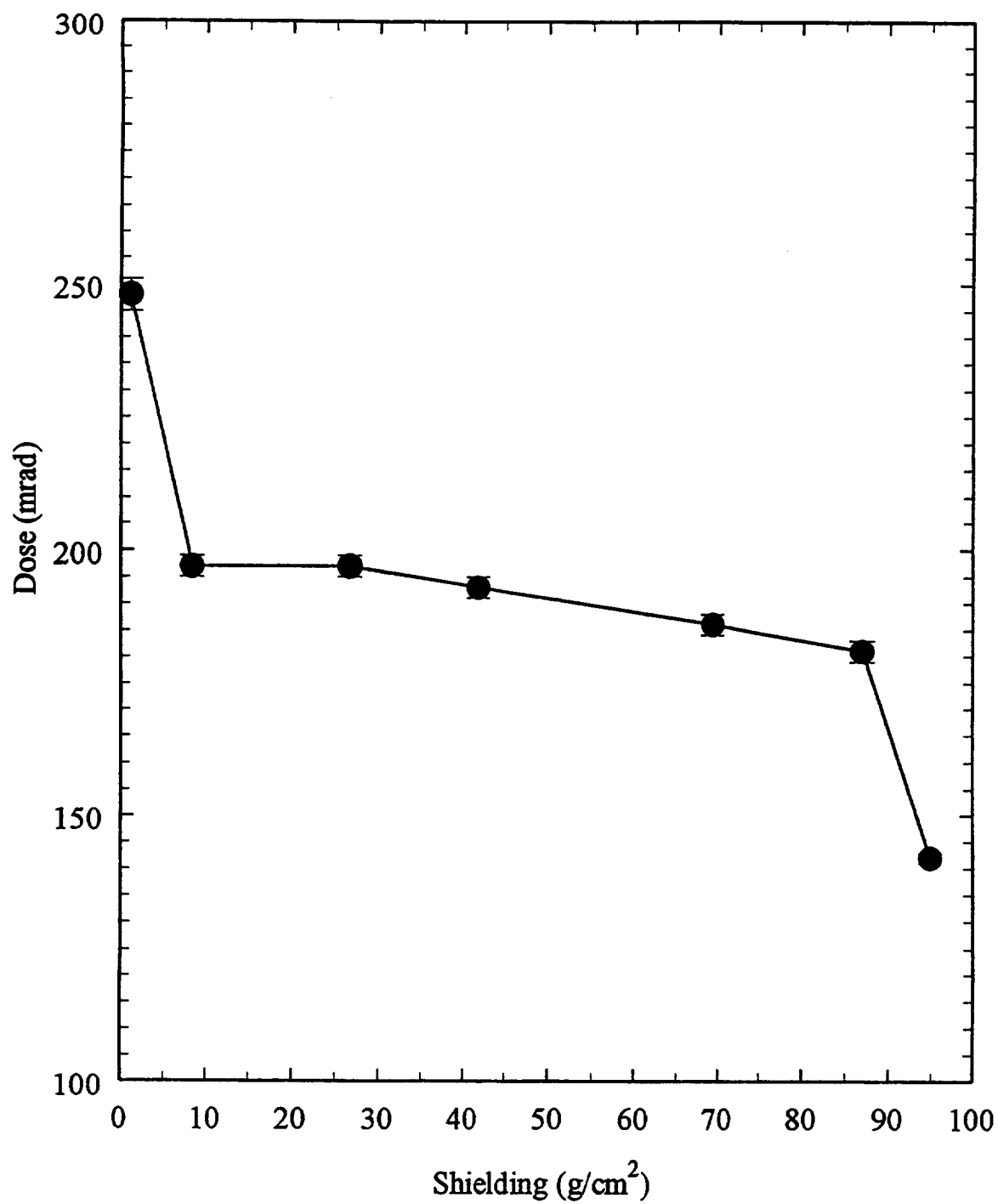


Figure 15: Mean TLD dose as a function of shielding depth from top to bottom in the ECT experiment stack.

container. Given the higher doses at mid-level values of shielding in TLD 5 versus the other locations, this location may have been under the minimum of shielding from surrounding experiments and from the orbiter itself. TLD Position 12 also shows an increase in dose between layers E19/E20 and E12/E05, illustrating a similar minimum of vertical shielding.

It is also possible that TLD Position 5 was facing toward the West, the direction of greatest trapped proton flux, while the spacecraft traversed the SAA, during the majority of the mission, leading to a higher dose from trapped protons in this particular TLD position. Since the spacecraft was oriented upside down with the open cargo bay facing the Earth, the vertical sides of the experiment were perpendicular to the Earth's surface during most of the mission. Trapped protons, travelling in a helical motion around the lines of the geomagnetic field tend to encounter the spacecraft parallel to the Earth's surface. Due to the East/West trapped proton anisotropy, the greatest flux of trapped protons impinges on the spacecraft from the West. STS-62 was an Earth observation mission and it is likely that the orientation of the orbiter remained fixed relative to the Earth during much of the mission. This would permit the East/West effect to be seen in the relative doses of TLDs positioned across the detector surface and may explain the substantially larger doses seen in TLD position 5 and, to a lesser degree, TLD position 12 as compared to doses measured in the other TLD positions.

5 Low Energy Neutron Fluences and Dose Equivalents

The thermal (<0.2 eV) and resonance (0.2 eV– 1 MeV) neutron fluences and dose equivalents were measured in the ECT assembly on STS-62 with Thermal/Resonance Neutron Detectors (TRNDs). The TRNDs are composed of ^6LiF films between CR-39 PNTD layers. Each detector contains a pair of $^6\text{LiF}/\text{CR-39}$ sandwiches with one of the pair covered by Gd foil ($25\text{ }\mu\text{m}$) thermal neutron absorbers. When irradiated with neutrons, the $^6\text{Li}(n,\alpha)\text{T}$ reaction takes place and the CR-39 layers are irradiated in turn by lower energy α -particles. In the half of the detector covered by Gd, the thermal neutrons are eliminated from the reactions, allowing a separation between thermal- and resonance-induced track densities on the CR-39. By assuming energy spectra for the incident neutrons (a well-thermalized spectrum below 0.2 eV; a E_n^{-1} moderated spectrum between 0.2 eV and 1 MeV) calibrations in neutrons/track

have been determined for the TRND.

The ECT experiment contained three TRNDs: Detector C (1 and 2) were placed at corners of the assembly below the chamber lid (under 0.306 g/cm^2), Detector B was centered in the lower calorimeter (under 44.34 and 50.57 g/cm^2 to top and bottom, respectively), Detector A was centered in the lower calorimeter (under 89.57 and 5.34 g/cm^2). A fourth detector (D) was shipped with the flight detectors in a Ground Control unit and provided background measurements.

After postflight return of the dosimeters, the TRNDs were disassembled and the CR-39 layers were processed for 4.5 hour in 6.25 N NaOH solution at 70°C . The α -particle track densities on the CR-39 were counted under $200\times$. The measured track densities are given in Table 5. Each track density is an average from the CR-39 layers above and below the ^6LiF /CR-39 film and is a measure of 4π neutron incidence (an average is used because TRNDs have been calibrated for a single CR-39 layer in contact with the ^6LiF film).

There is a large variation in track densities in Table 5 with Detector C only about 3 times greater than the Ground Control (D). The most centered position (B) has track densities ~ 10 times greater than Detector C and 3 times greater than Detector A. Not shown is the fact that in Detector A (near the bottom of the assembly) the track densities from the top CR-39 layer were ~ 3 times larger than those on the bottom CR-39 layer. Low energy neutron densities clearly fall off sharply near the edges of the ECT assembly. The same difference was not seen between top and bottom CR-39 layers in Detector C (near the top corners of the assembly) where all track densities were low.

Track densities are converted to neutron fluences and dose equivalents in Table 6. Detector calibration has been described by Benton et al.[1]. Dose equivalent conversion factors from NCRP (1971) were used which incorporated QF values of 2 for thermal neutrons and 6.4 for resonance neutrons. The low energy neutron dose equivalents of 11.6, 39, and 3.1 mrem (of 1.03, 3.5 and 0.28 mrem/d) can be compared with previous Shuttle measurements. The average of 11 flight measurements (from a variety of altitudes and inclinations) is 0.26 mrem/d with a σ of 0.12 mrem/d. The dose equivalent in the interior of the ECT assembly is much higher than previous measurements and demonstrates that substantial neutron moderation and thermalization takes place within a hydrogenous mass. The TLD

Table 5: Track Densities from the ECT TRNDs

Detector	Total Track Density (cm^{-2})	Background Subtracted Track Density (cm^{-2})
A	1311 ± 42	1223 ± 43
A-Gd	439 ± 18	394 ± 19
B	3597 ± 87	3509 ± 87
B-Gd	1370 ± 40	1325 ± 40
C	235 ± 12	147 ± 14
C-Gd	152 ± 10	107 ± 12
D	88 ± 7	
D-Gd	45 ± 6	

Gd designates detectors with Gadolinium covers.

Detector D was a Ground Control unit and the track densities represent background for the flight detectors.

Positions of the A, B, and C detectors were at the top, middle and bottom, respectively, of the ECT assembly.

absorbed dose near Detector B was about 170 mrad and lower energy neutron dose equivalent represents less than 23% of the total.

6 Conclusion

Measurements of LET spectra were fairly typical for a STS mission of this altitude and inclination. STS-62 was in a 39° , 296 km orbit, falling between the minimum and maximum orbital inclinations for the Space Shuttle of 28.5° and 57° , respectively. Thus STS-62 received significant fractions of its radiation exposure from the ionizing radiation components that dominate these two extremes: stopping primary protons and short range, high LET secondaries from trapped protons in the SAA for low inclination orbits and GCR for high inclination orbits. Little difference was seen amongst the seven measured integral LET flux spectra as a function of shielding. The largest flux in the low LET region ($<10 \text{ keV}/\mu\text{m}$)

Table 6: Neutron fluence and Dose Equivalent from ECT TRNDs

Detector	Energy Range	Neutron Fluence (cm^{-2})	Dose Equivalent (mrem) (μSv)	
A	<0.2 eV	$3.0 \pm 0.6 \times 10^5$	0.30 ± 0.06	3.0 ± 0.6
	0.2 eV–1 MeV	$2.3 \pm 1.1 \times 10^6$	11.3 ± 5.6	113 ± 56
B	<0.2 eV	$7.8 \pm 1.6 \times 10^5$	0.79 ± 0.16	7.9 ± 1.6
	0.2 eV–1 MeV	$7.7 \pm 3.8 \times 10^6$	38 ± 19	380 ± 190
C	<0.2 eV	$1.4 \pm 0.7 \times 10^4$	0.015 ± 0.007	0.15 ± 0.07
	0.2 eV–1 MeV	$6.2 \pm 3.1 \times 10^5$	3.1 ± 1.5	31 ± 15

was seen in detector pair E33/E25 under 1.19 g/cm^2 (Figure 6) and most likely took the form of stopping primary protons encountered in the SAA. The largest fluxes in the mid ($>10 \text{ keV}/\mu\text{m}$) and high ($>100 \text{ keV}/\mu\text{m}$) regions were seen in detector pairs E15/E14 (Figure 8) and E19/E20 (Figure 9), and were probably mainly secondaries produced by trapped protons in the SAA. Previous work on the Long Duration Exposure Facility (LDEF) demonstrated that the production of proton-induced secondaries increases with shielding[2]. While E15/E14 and E19/E20 were in the middle range of shielding in the experiment (26.71 and 41.84 g/cm^2 , respectively) they may have been the most heavily shielded layers when the additional shielding of the spacecraft and surrounding experiments is taken into account and it is this total shielding that is responsible for the large number of proton-induced secondary particle tracks seen in these detectors.

Approximately 20% of the mid to high LET particle events were long range ($>600 \mu\text{m}$) and thus can be considered to have been produced by GCR. Of the long range particles, 80% were slowing; that is a noticeable difference in size could be seen between tracks on each of the two detector surfaces. This permitted the direction of these particles to be determined and it was found that the large majority of long range particles arrived in the experiment stack from the direction opposite the Earth and passed through the stack towards the Earth. This illustrates the effect of the large amount of shielding ($\sim 100 \text{ g/cm}^2$) of the ECT experiment

on GCR.

Total absorbed dose as measured by TLDs was seen to be attenuated by shielding in the experiment stack. The least shielded layer (1.19 g/cm^2) had a mean dose of $248 \pm 3 \text{ mrad}$. Dose decreased rapidly in the first $\sim 7 \text{ g/cm}^2$ of shielding and then plateaued at $\sim 190 \text{ mrad}$ in the middle of the experiment stack. At the bottom of the stack, under the greatest shielding of 94.91 g/cm^2 , mean dose dropped to $142 \pm 1 \text{ mrad}$. Little difference in dose could be seen for the nine TLD locations across a given detector layer, except for TLD Position 5. This position measured a significantly higher dose in the middle levels of the experiment stack and was on the left-most edge of the detector stack. This difference in measured dose as a function of TLD position was probably due to a combination of shielding and experiment orientation. Previous work on LDEF[3] demonstrated the differences in dose across a spacecraft orbiting in a fixed orientation relative to the Earth due to the East/West trapped proton anisotropy in the SAA. Since STS-62 was an Earth observation mission, it is reasonable to assume that the orbiter remained in a fixed orientation for a significant portion of the mission. It is possible that TLD Position 5 was oriented toward the West and under comparatively low shielding, leading to a disproportionately large dose from trapped protons at this position as compared to other TLD positions across the detector surface.

Thermal and resonance neutron dose equivalents were found to vary substantially with shielding. In this case shielding can be considered a neutron thermalizing and moderating medium, since it was largely hydrogenous in composition. A low energy dose equivalent of 3.5 mrem/d was measured in Detector B under 41.84 g/cm^2 . This is substantially greater than the 0.26 mrem/d mean dose equivalent measured on previous STS missions, demonstrating the effect of the hydrogenous mass of the experiment in neutron moderation and thermalization.

Since the ECT experiment was included on an Earth observation STS mission and thus orbited upside down during most of the flight, the primary objectives of the experiment could not be met. However, the fact that a large percentage (80%) of long range particle events were found to be traversing the experiment from space towards the Earth and were seen to be slowing down validates the general design of the experiment. LET spectra measurements showed that a substantial fraction (80%) of mid to high LET particles took the form of short

range, high LET secondaries produced by trapped protons and indicates that this component is significant in orbits of this inclination (39°) as well as lower inclination orbits and must be adequately taken into account during the data reduction and analysis process.

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APPENDIX B

ECT Mechanical, Thermal, and Electrical Systems Report

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1.0 Summary

This report documents the preparation and performance of the Emulsion Chamber Technology Experiment (ECT) hardware and software for the OAST-2/STS-62 mission aboard the Orbiter Columbia. Also discussed are lessons learned, current status and issues for future flights of ECT.

ECT flew as one of several experiments comprising the OAST-2 payload on STS-62, beginning with launch at KSC on 4 March 1994 and ending with touchdown at KSC on 18 March 1994. The ECT flight hardware was located on the upper surface of the Hitchhiker-M bridge structure in the aft end of the payload bay, next to the portside payload bay sill. The hardware was facing out of the payload bay in the +Z direction, affording an acceptable view. The effective exposure time accrued was seventy-five (75) hours. Columbia's flight performance was nominal and impacts on experiment operations were minimal.

Excellent thermal control of the ECT Emulsion Chamber was maintained during flight and no significant power interruptions occurred. Telemetry and recovered on-board data confirm that the temperature of the Emulsion Chamber was controlled to 20 ± 1 °C throughout the entire flight. The flight hardware arrived back at MSFC after the OAST-2/STS-62 mission in good condition. The emulsion stack showed no apparent thermal, mechanical, light or radiation damage and contained returned data. All stored electronic data was recovered.

The mechanical interference problem identified during system-level integration and testing at GSFC involving the pressure transducer and the adjacent mounting bolt was eliminated by correct positioning of the pressure transducer during final assembly of the Emulsion Chamber. No interference problems were encountered during pre-launch integration at KSC. None of the thermal switches de-bonded during shipment, ground handling or flight.

2.0 Current Status

The flight hardware, quality-sensitive spares and quality-sensitive ground support equipment (GSE) are packed for storage in the ECT Shipping Container. All AA alkaline cells have been removed from the Electronics Unit and the Temp Mentor. Quality Assurance documentation and the Hardware Activity Log are packed with the hardware.

Quality Assurance coverage has been maintained by NASA since hardware delivery and acceptance. MSFC Quality Assurance personnel have indicated that QA coverage will be continuous and that ECT will be ready for re-flight provided that all quality-sensitive items are placed in appropriate storage at MSFC. As of the time of this report, the flight

hardware, quality-sensitive spares, quality-sensitive GSE and non-quality-sensitive items have been placed in bonded storage at MSFC.

Emulsion stack materials, including the lead sheets and processed plates, are in the custody of the Cosmic Ray Emulsion Laboratory (MSFC/ES62). Development hardware and other non-quality-sensitive or non-flight items are currently stored at UAH Materials Science Building. Post flight data products provided by NASA will also be found at UAH Materials Science Building.

3.0 Assembly, Integration and Flight

3.1 Assembly for Flight

Previous reports have described the system-level integration and testing at GSFC. The Temp Mentor's RS-232 power loop-back feature caused the Temp Mentor on-board power depletion while at GSFC. Thus, no temperature record for the period 10/26/93 through 11/23/93 exists. A mechanical interference problem was identified involving the pressure transducer and the adjacent mounting bolt. The bolt was omitted for GSFC testing. The ECT hardware was de-integrated and shipped back to MSFC from GSFC after completion of the required integration and testing activities, arriving on 11/23/93. The test emulsion stack arrived with no apparent damage. The ECT hardware also arrived in excellent condition. The materials for the flight emulsion stack had been made ready by this time at the Cosmic Ray Emulsion Laboratory (MSFC/ES62).

3.1.1 Emulsion Chamber Assembly

The Emulsion Chamber was cleaned thoroughly before final assembly for flight. The loading of the flight emulsion stack went extremely well once again. The assembly of the Emulsion Chamber with emulsion stack has now been successfully completed three times, once with the development hardware and twice with the flight hardware. The use of alignment fixtures during assembly was abandoned early on and the standard procedure is to assemble the Bottom Plate and Lower Chamber before loading. The Emulsion Chamber assembly procedure is included in Appendix A. The flight emulsion stack configuration record was provided by the Cosmic Ray Emulsion Laboratory (MSFC/ES62) and is included in Appendix B.

The pressure transducer was rotated to the extreme counter-clockwise or (nearly) vertical position when installed on the Emulsion Chamber, in order to alleviate the interference problem involving the pressure transducer and the adjacent mounting bolt. As a result, no mechanical interference was encountered during integration of the ECT flight hardware at KSC. This step of the Emulsion Chamber assembly must be performed with

the ECT Flight Wiring Harness connected to the Emulsion Chamber in order to ensure proper positioning of the pressure transducer while preventing interference with the adjacent feedthrough connector.

The O-rings used in the assembly of the Emulsion Chamber were coated with a minimum amount of vacuum grease. Leak testing followed assembly, with Helium detection and pressure decay tests. In keeping with the ECT mission approach, no extraordinary methods were used. It was noted during testing that considerable amounts of Helium had been absorbed by the stack materials, evidenced by unexpected pressure increases during pressure decay tests. The Emulsion Chamber was left pressurized with air to approximately 30 psia for shipment to KSC.

3.1.2 Electronics Unit Assembly

The Electronics Unit was cleaned thoroughly before final assembly for flight. Fresh AA alkaline cells were installed in the Electronics Unit and the Temp Mentor. A flight-qualified PROM hosting the 7/93 version of the ECT flight software was installed in the Electronics Unit. This version of the flight software was verified by over 150 hours of test operation before installation in the Electronics Unit.

The power loop-back feature was disabled on the Temp Mentor's RS-232 port and instead built into the Electronics Unit's RS-232 port, feedthrough connector J7. Power is now conserved because GSE cables are connected to the Electronics Unit's RS-232 port for only a relatively short time, during ground operations.

3.2 Final Preparations for Shipment

Two bimetallic thermal switches were re-bonded to the Emulsion Chamber with flight epoxy. Flight velcro was installed on both assemblies, lockwires were installed, lockwire ends were encapsulated with flight epoxy, and touch-ups were made on heater wire insulation and silverized Teflon tape. After some assurances were given by GSFC concerning ferry temperature control, the decision was made to omit the Solimide TA-301 polyimide foam insulation, due to its tendency to shed particulates. It is likely that a method could be developed using adhesive tape or a hot surface to seal the surfaces and make the foam acceptable for flight. The concerns raised last summer by GSFC's QA contractor and others were addressed with these actions.

All exposed connectors on the Emulsion Chamber and Electronics Unit were covered with Teflon dust covers before shipment. The Temp Mentor was deployed and the ECT Shipping Container was sealed for shipment on 12/29/93. Shipment to KSC occurred on 1/3/94.

3.3 Integration for Flight

The ECT team arrived at KSC on 1/4/94. OAST-2 integration and testing activities took place in Hangar AM. The hardware was unpacked, checked out and installed on the Small Top Pallet. The Emulsion Chamber was vented to atmosphere. The Temp Mentor data was recovered and the Temp Mentor was deployed for flight. The Small Top Pallet with the ECT flight hardware was then installed on the HH-M structure. Power and data cables were integrated, followed by installation of the MLI and ground wires. The Emulsion Chamber was again vented to atmosphere just before the MLI was closed. GSFC provided some grounding lugs, lockwires and cable ties, cleaning of the radiator surfaces and some stitching on the MLI. Integration and testing activities were completed on 1/6/94. OAST-2 was integrated with the Orbiter Columbia some weeks later.

3.4 The OAST-2 Mission

ECT flew as one of several experiments comprising the OAST-2 payload on STS-62, beginning with launch at KSC on 4 March 1994 and ending with touchdown at KSC on 18 March 1994. The ECT flight hardware was located on the upper surface of the Hitchhiker-M bridge structure in the aft end of the payload bay, next to the portside payload bay sill. The hardware was facing out of the payload bay in the +Z direction, affording an acceptable view. The effective exposure time accrued was seventy-five (75) hours. Columbia's flight performance was nominal and impacts on experiment operations were minimal.

The ECT team staffed the GSFC Payload Operations Control Center (POCC) around the clock during the OAST-2/STS-62 flight. Training sessions and mission simulations were also supported in the weeks prior to the mission.

3.5 De-integration

The ECT flight hardware was de-integrated from OAST-2 and packed for shipment to MSFC on 5 April 1994 at KSC. The ECT Shipping Container, along with spares and materials arrived at MSFC early on 6 April 1994. Disassembly, processing of emulsion stack materials and packing for storage followed. The Hardware Activity Log is included in Appendix C.

4.0 Engineering Performance

4.1 Thermal Performance

The performance of the ECT thermal control system was excellent. The OAST-2 payload experienced a cold soak period after reaching orbit, due to Orbiter operations before OAST-2 power-up. Early telemetry indicated that the temperature of the emulsion stack ranged from approximately 9 °C (after correction) at the top of the stack (near the space radiator) to approximately 16 °C at the middle and bottom locations. Power was applied to OAST-2 and its experiments, including ECT, at MET 00:02:39 (Mission Elapsed Time, dd:hh:mm), or two hours, thirty-nine minutes after launch. Upon power-up the ECT thermal control system began operation and Emulsion Chamber temperatures began to recover. At no time did the emulsion stack temperature fall below 5 °C. Figures 1 and 2 provide the internal temperatures and power consumption for ECT following power-up. From the data contained in Figure 1, the thermal time constant of the emulsion stack is estimated to be approximately five (5) hours.

After recovery from the cold soak, the temperature of the ECT Emulsion Chamber was maintained at 20 ± 1 °C in all spacecraft attitudes throughout the entire flight. Orbiter performance was nominal and no significant interruptions of experiment power occurred. Telemetry and recovered on-board data confirm that the temperature of the Emulsion Chamber was controlled to 20 ± 1 °C. Correction of the readings for the emulsion stack top thermistor (Appendix E) is required as described in the thermistor calibration report contained in Appendix D. After correction, these values do indeed fall within the specified range of 20 ± 1 °C.

The temperature of the Small Top Pallet remained within the range of 0 - 40 °C in all spacecraft attitudes throughout the entire flight, as promised by GSFC. It was reported that the Small Top Pallet heater was never needed during the flight.

All of the telemetry data relevant to ECT flight operations is presented in Appendices E - I. These include the three emulsion stack temperatures, ECT power and the Small Top Pallet temperature, recorded at thirty-minute intervals at the POCC at GSFC.

Previous reports have presented analytical predictions and thermal test results. Predictions, test results and flight results are shown in Table 1. In all analyses and tests Emulsion Chamber temperatures were controlled to within the range of 20 ± 1 °C unless the analysis or test reports indicated otherwise.

ECT power consumption varies considerably with Orbiter attitude. Generally, predictions and test environments were conservative as compared to the flight data.

Figure 1. ECT Internal Thermistors, MET 0/02:40:00 - MET 0/04:10:00, Power-up

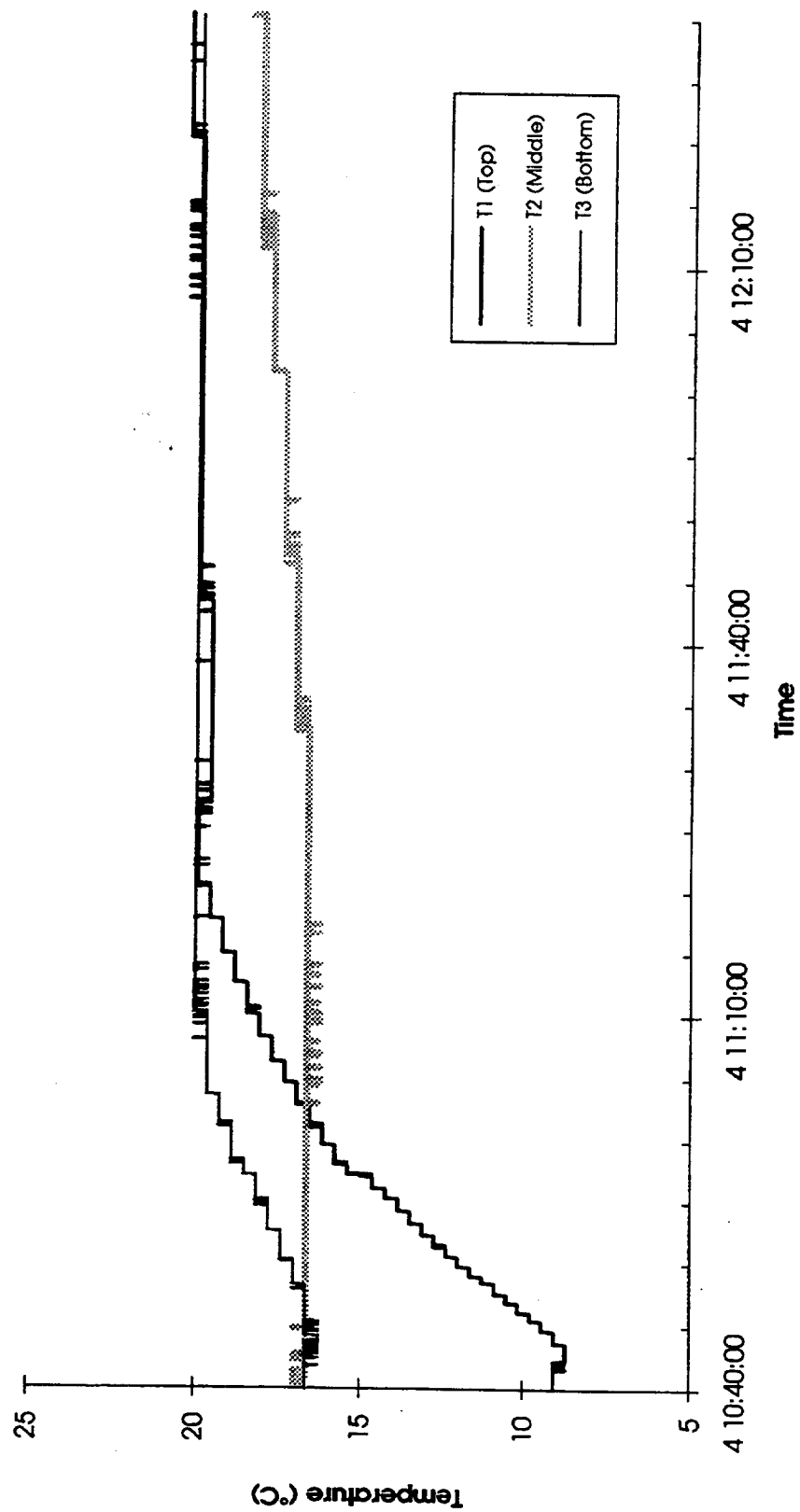


Figure 2. ECT Power Draw, MET 0/02:40:00 - MET 0/04:10:00, Power-up

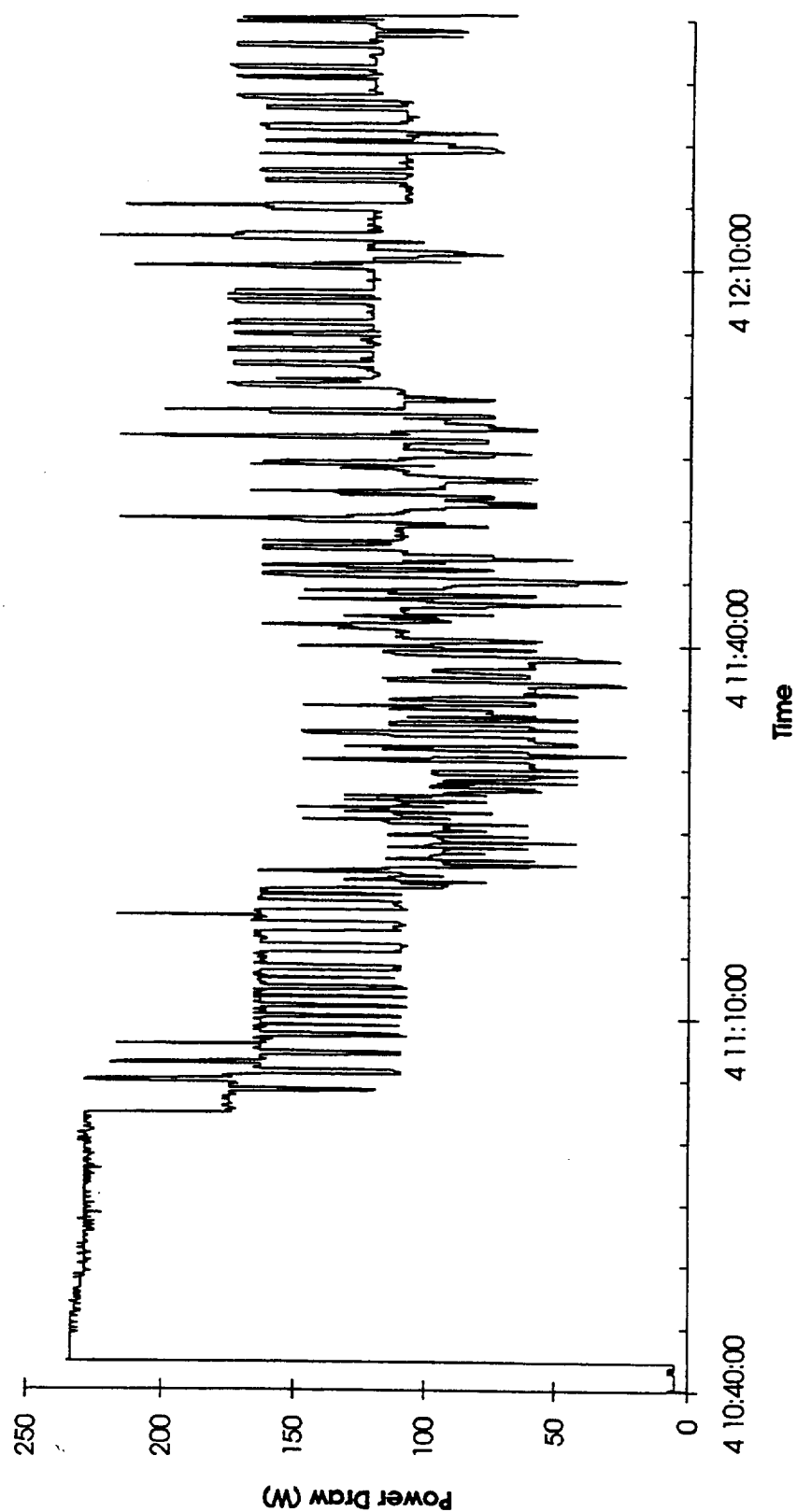


Table 1. Comparison of Predictions, Test Results and Flight Results

Orbiter Attitude	Predicted Avg. Power (W)	Tested Avg. Power (W)	Flight Avg. Power (W)
Bay-to-Sun	38	76	61
Tail-to-Earth	not predicted	not tested	95
Bay-to Earth	78	76	54
Bay-to-Space	112	131	109

4.1.1 Bay-to-Sun Attitude

The predicted power value was expected to be rather low compared to test and flight results. This is due to the fact that the bay-to-sun thermal environment was supplied by the GSFC thermal contractor for OAST-2's first flight assignment and was not updated (because it was such a conservative environment) until after the UAH thermal analysis was completed, but before thermal testing was conducted.

This attitude was used a few times for thermal conditioning of the payload bay before extended periods in cold attitudes. The period MET 09:18:08 - MET 09:19:08 is the best and longest example. Temperature and power data for the period MET 09:18:08 - MET 09:19:08 are shown in Figures 3 and 4.

4.1.2 Tail-to-Earth Attitude

This attitude was flown to accommodate OAST-2. The period MET 09:19:08 - MET 09:20:28 is one example. Temperature and power data for the period MET 09:19:08 - MET 09:20:28 are shown in Figures 3 and 5. No thermal environment was provided before the mission.

4.1.3 Bay-to Earth Attitude

This attitude was flown for the bulk of the mission to accommodate the primary payload. Temperature and power data for the period MET 03:03:21 - MET 03:05:02 are shown in Figures 6 and 7.

4.1.4 Bay-to-Space Attitude

This attitude was flown a few times to accommodate OAST-2 and certain Orbiter operations. The period MET 10:06:51 - MET 10:08:36 is one example. Temperature and power data for the period MET 10:06:51 - MET 10:08:36 are shown in Figures 8 and 9.

Figure 3. ECT Internal Thermistors, MET 9/18:08 - MET 9/20:28, Bay-to-Sun & Tail-to-Earth

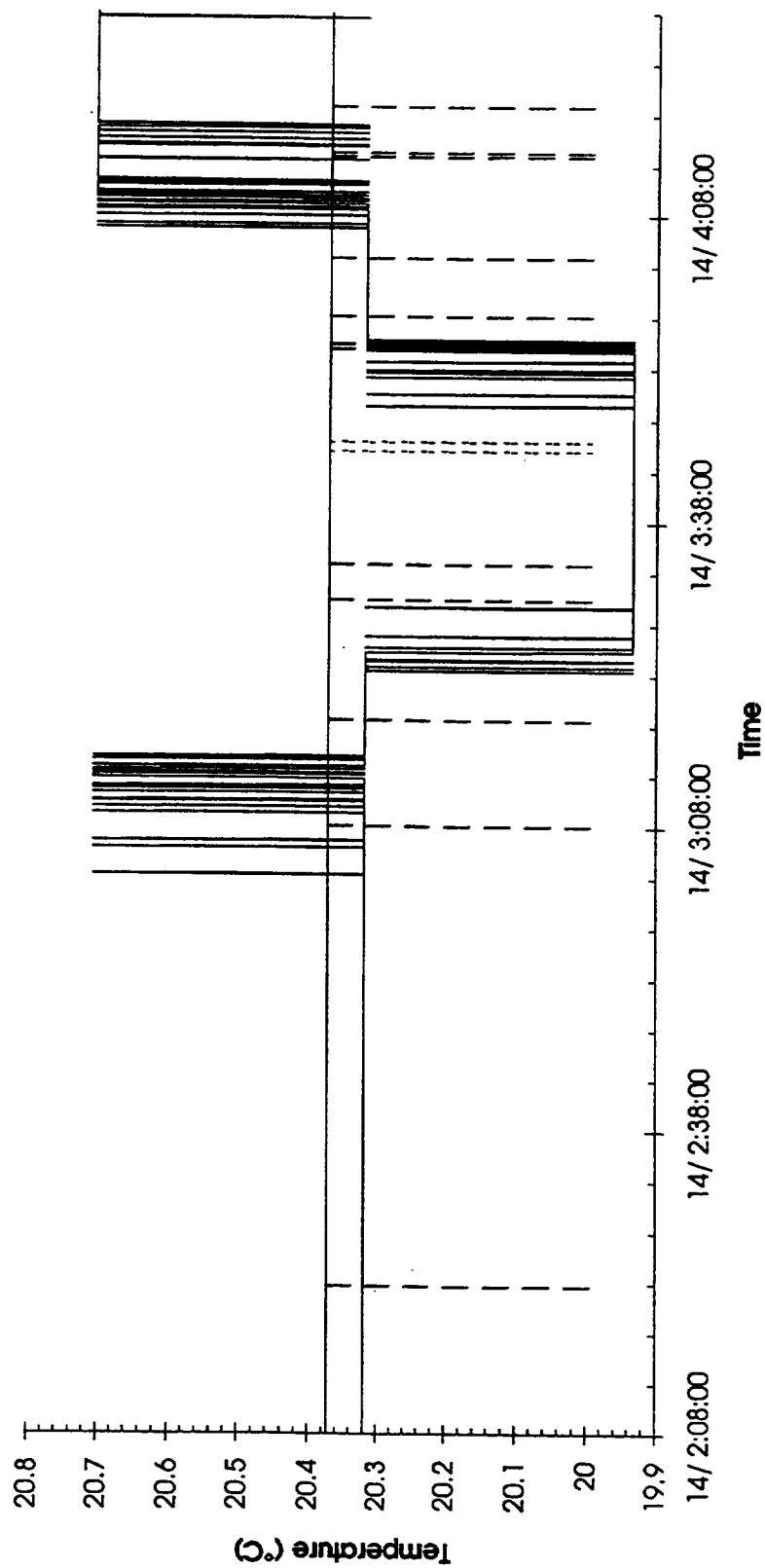
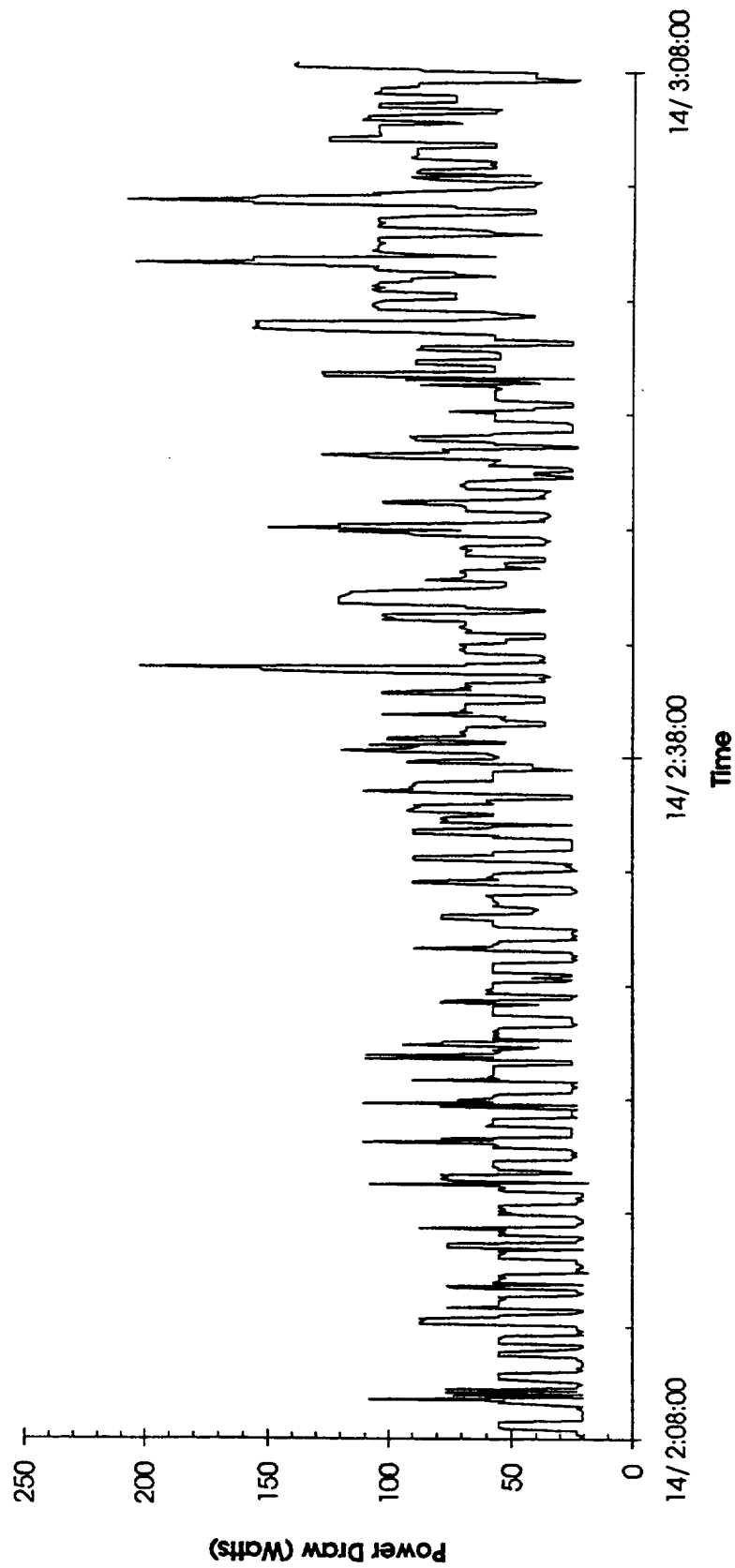
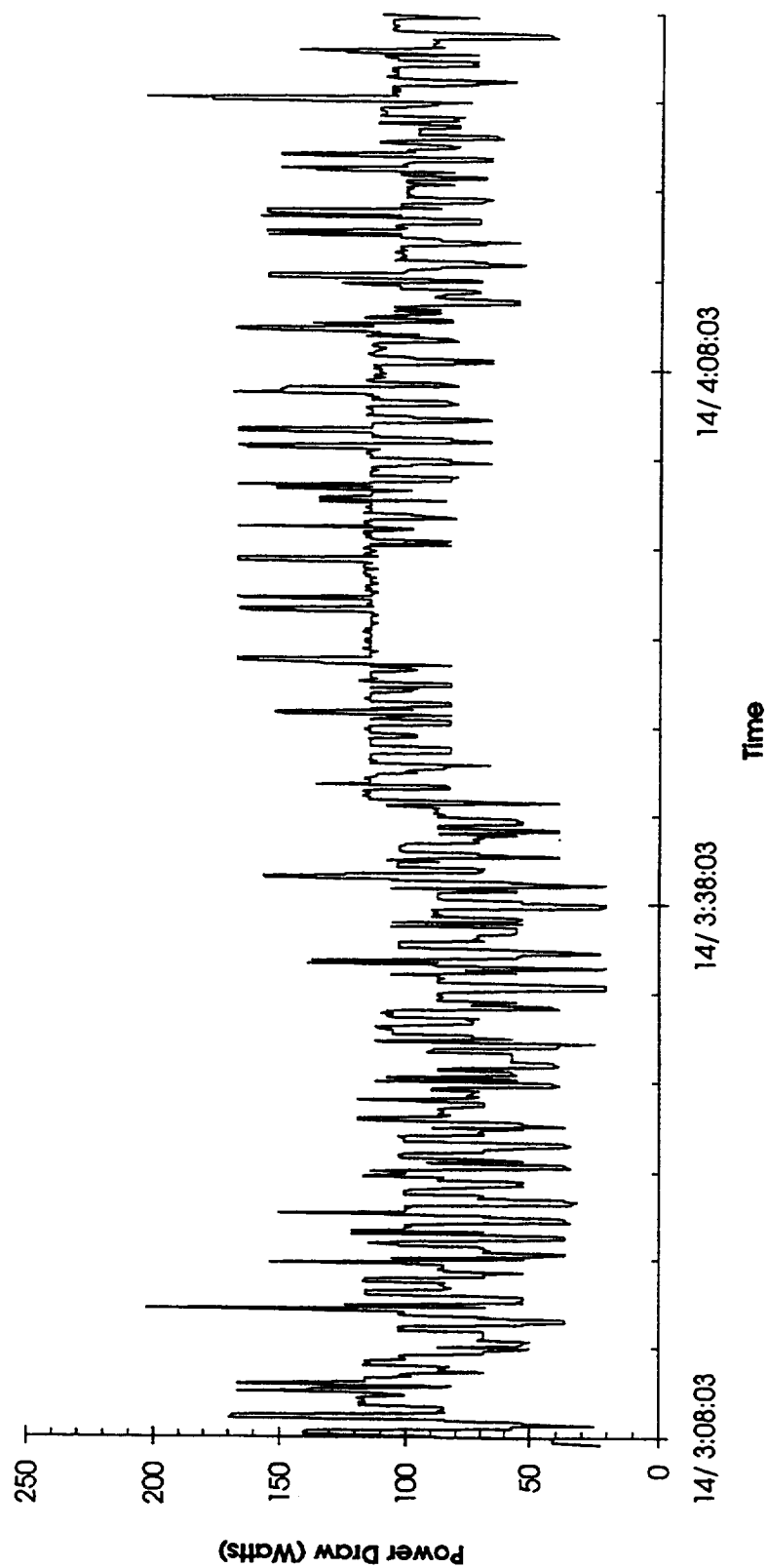


Figure 4. ECT Power Draw, MET 9:18:08 - MET 9:19:08, Bay-to-Sun



Bay-to-Sun Avg. Pwr. 61W

Figure 5. ECT Power Draw, MET 9:19:08 - MET 9:20:28, Tail-to-Earth



Tail-to-Earth Avg. Pwr. 95W

Figure 6. ECT Internal Thermistors, MET 3/03:21 - MET 3/05:02, Bay-to-Earth

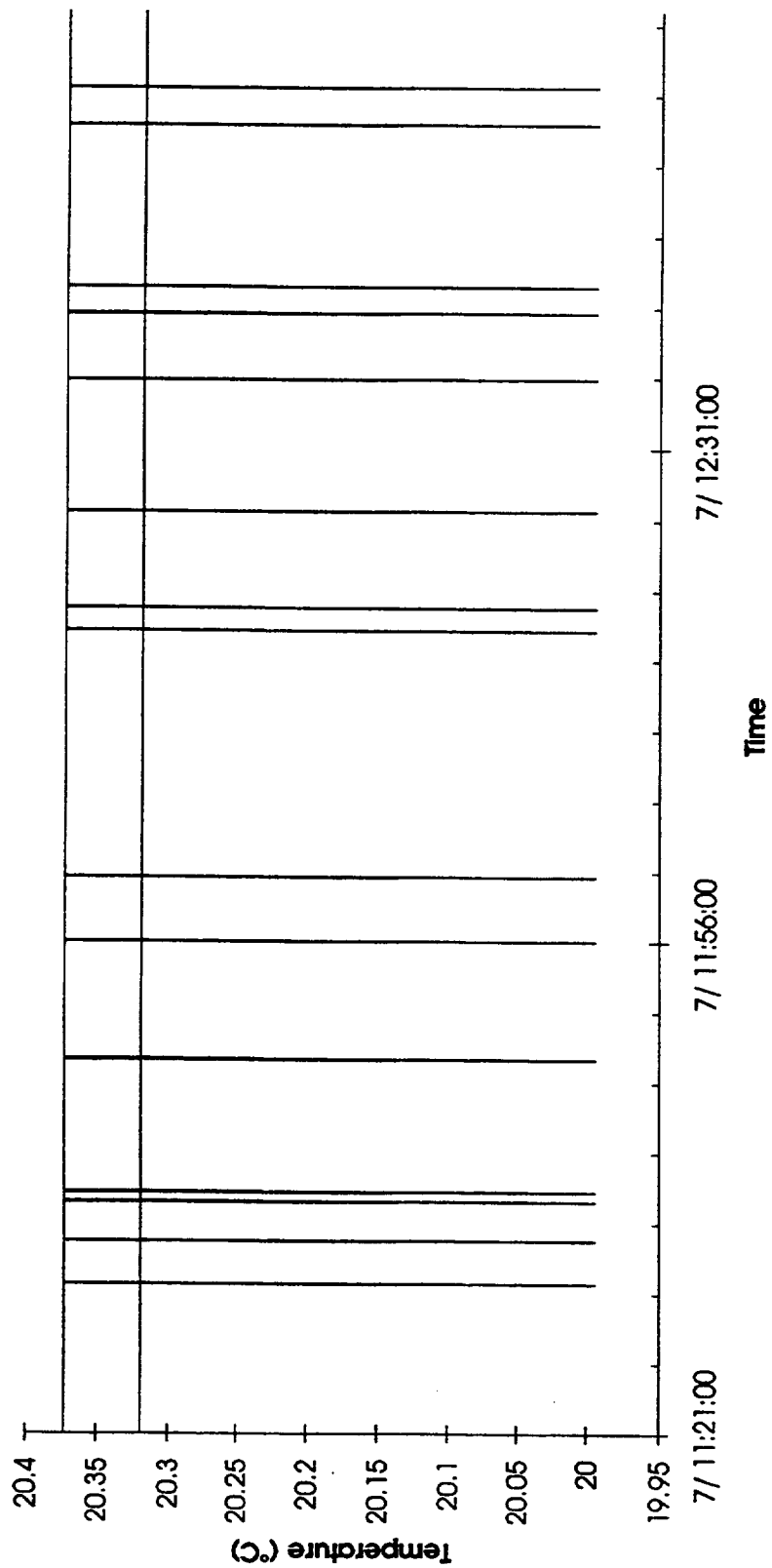
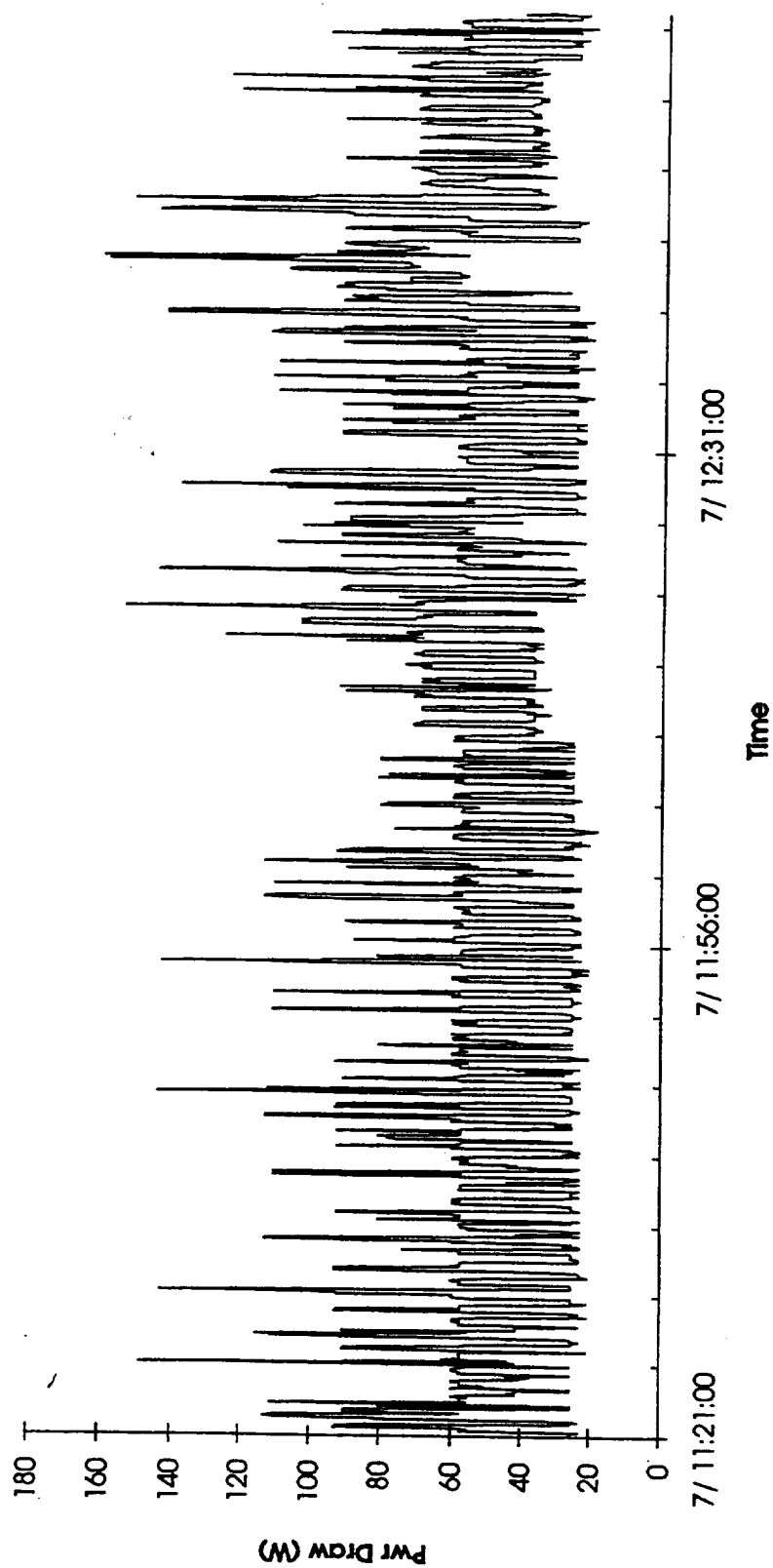


Figure 7. ECT Power Draw, MET 3/03:21 - MET 3/05:02, Bay-to-Earth



Bay-to-Earth Avg. Pwr. 54W

Figure 8. ECT Internal Thermistors, MET 10/06:51 - MET 10/08:36, Bay-to-Space

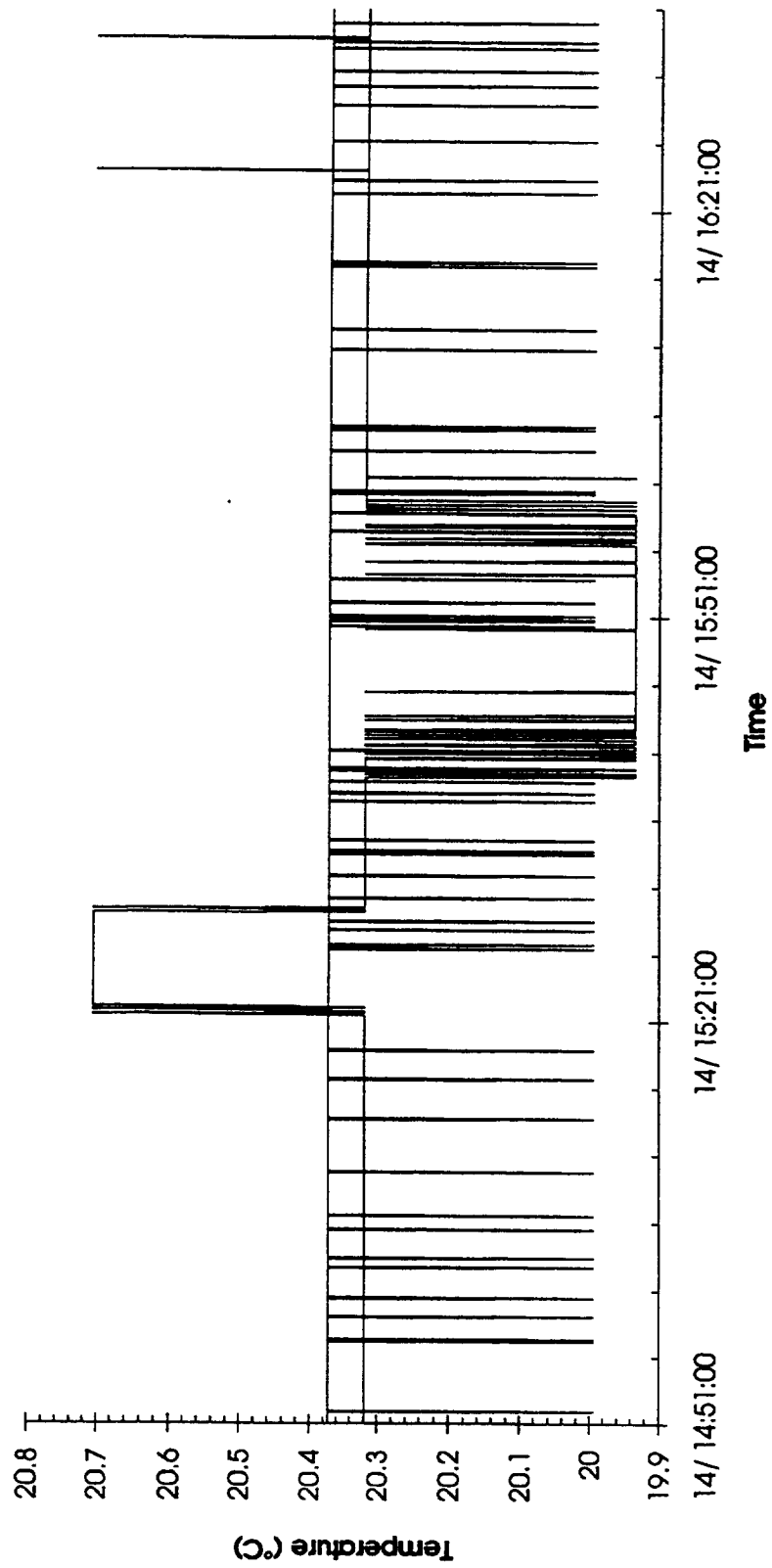
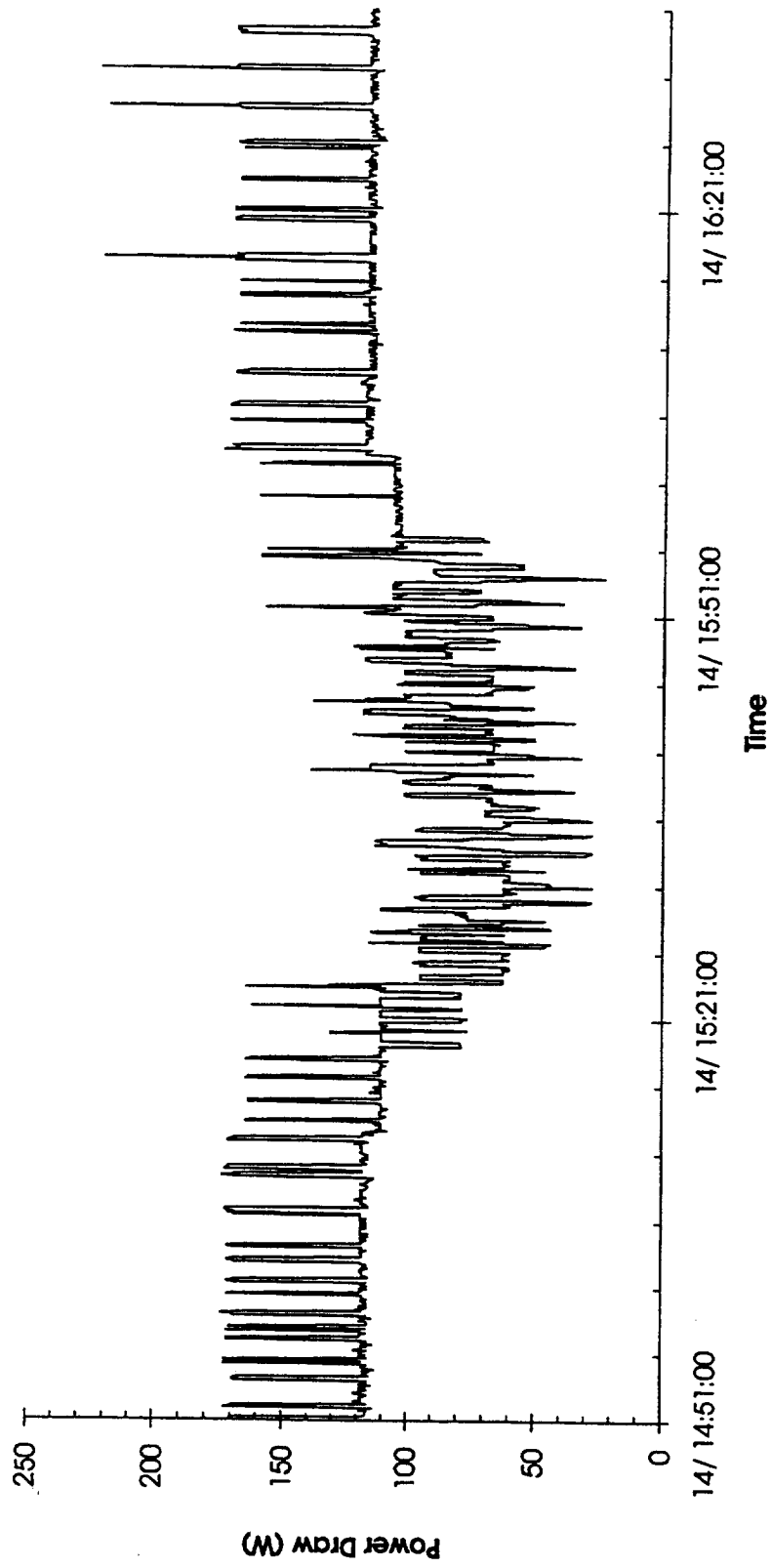


Figure 9. **ECT Power Draw, MET 10/06:51 - MET 10/08:36, Bay-to-Space**



Bay-to-Space Avg. Pwr. 109W

4.2 Mechanical Performance

The ECT flight hardware and GSE mechanical performance was nominal. No damage or unusual wear was noted and all flight parts, including fasteners, were recovered after de-integration and disassembly. No mechanical spares were required for the OAST-2/STS-62 mission. No mechanical damage to the flight emulsion stack or evidence of light leakage has been reported to date.

4.2.1 Mechanical Interference

A mechanical interference problem was identified during GSFC integration and testing involving the pressure transducer and the adjacent mounting bolt. Gerard Durback (GSFC) elected to omit the mounting bolt during GSFC integration and testing. Upon final assembly, the pressure transducer was rotated to the extreme counter-clockwise or (nearly) vertical position when installed on the Emulsion Chamber, in order to alleviate this interference problem. No mechanical interference was encountered during integration of the ECT flight hardware at KSC and all mounting hardware was installed for flight.

4.2.2 Mechanical Installation

Installation of the Emulsion Chamber onto the Small Top Pallet for flight was performed by GSFC personnel with assistance from the ECT team and mounting hardware torques and gaps were set according to the procedure agreed to by Gerard Durback (GSFC) and Dr. Fran Wessling (UAH). All thermal insulation hardware was installed. During installation it was determined that some mounting hardware gaps were present on the upper side of the Emulsion Chamber mounting flange and some were present on the lower side. Gerard Durback (GSFC) elected to employ shims in either or both positions such that the total of the two gaps would be within specification. Gerard Durback (GSFC) also elected to omit all shims on the Electronics Unit mounting hardware. Final mounting bolt torque values were 160 - 170 inch-pounds for both the Emulsion Chamber and the Electronics Unit. The GSFC installation procedure is included in Appendix J.

4.2.3 Emulsion Chamber Internal Pressure

As shown in Figure 10, the Emulsion Chamber maintained a steady internal pressure after some initial transients. In keeping with the ECT mission approach, no extraordinary methods were used in the fabrication or assembly of the Emulsion Chamber to ensure that internal pressure is maintained during orbital flight. The O-rings used in the assembly of the Emulsion Chamber were coated with a minimum amount of vacuum grease. Leak testing followed assembly, with Helium detection and pressure decay tests. It was noted during testing that considerable amounts of Helium had been absorbed by

the stack materials, evidenced by unexpected pressure increases during pressure decay tests. The Emulsion Chamber was left pressurized with air to approximately 30 psia for shipment to KSC.

4.2.4 Epoxy Adhesive Bonding of Thermal Switches

An effective procedure for bonding thermal switches to the flight hardware with epoxy adhesive was developed. The ECT epoxy adhesive bonding procedure is included in Appendix K. Surface preparation and cleanliness were found to be of the utmost importance. Extra washing of the bonding site with acetone seemed to be helpful and was usually included after the last cleaning step. Our recommendation for any future use of thermal switches is that the stud-mounted variety be employed if possible, preventing the need for bonding the parts.

4.3 Electrical & Data Performance

The Electronics Unit system operated nominally. Thermal control of the flight hardware was uninterrupted. The AA alkaline cells for the Electronics Unit on-board memory back-up power performed nominally and the stored record from the Electronics Unit on-board memory was recovered intact, if incomplete (refer to Section 4.3.2), after the flight hardware was returned to MSFC. See Figures 10 - 13.

Temp Mentor operation was nominal. The Temp Mentor on-board memory was more than adequate, the AA alkaline cells performed nominally and the stored record from the Temp Mentor on-board memory was recovered intact after the flight hardware was returned to MSFC. See Figures 14 and 15.

4.3.1 Temp Mentor

The Temp Mentor RS-232 port power loop-back feature energizes the Temp Mentor RS-232 interface whenever the Temp Mentor interface cable is plugged into the Temp Mentor RS-232 port, as it is when carried aboard the Electronics Unit. (This was the cause of the Temp Mentor power depletion while at GSFC. Thus, no temperature record for the period 10/26/93 through 11/23/93 exists.) This power loop-back was disabled on the Temp Mentor's RS-232 port and instead built into the Electronics Unit's RS-232 port, feedthrough connector J7, before delivery to KSC. Temp Mentor power is now conserved because GSE cables are connected to the Electronics Unit's RS-232 port, feedthrough connector J7, for only a relatively short time, during ground operations.

If a different Temp Mentor unit is intended for flight aboard the Electronics Unit in subsequent missions the same RS-232 port power loop-back must be disabled on that Temp Mentor's RS-232 port.

Figure 10. Emulsion Chamber Pressure

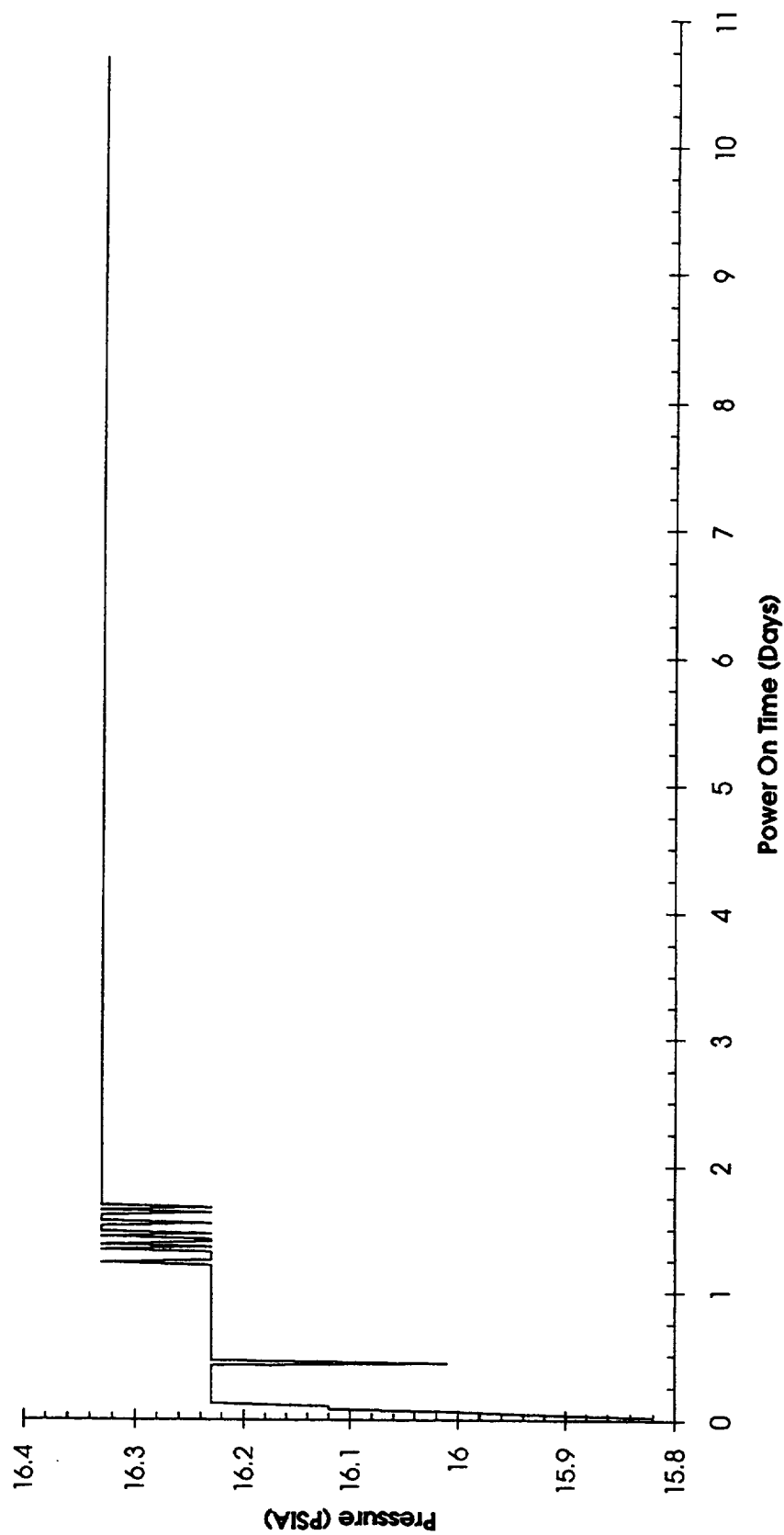


Figure 11. Chamber Top Thermistor Temperatures (Fine Channels 1-6)

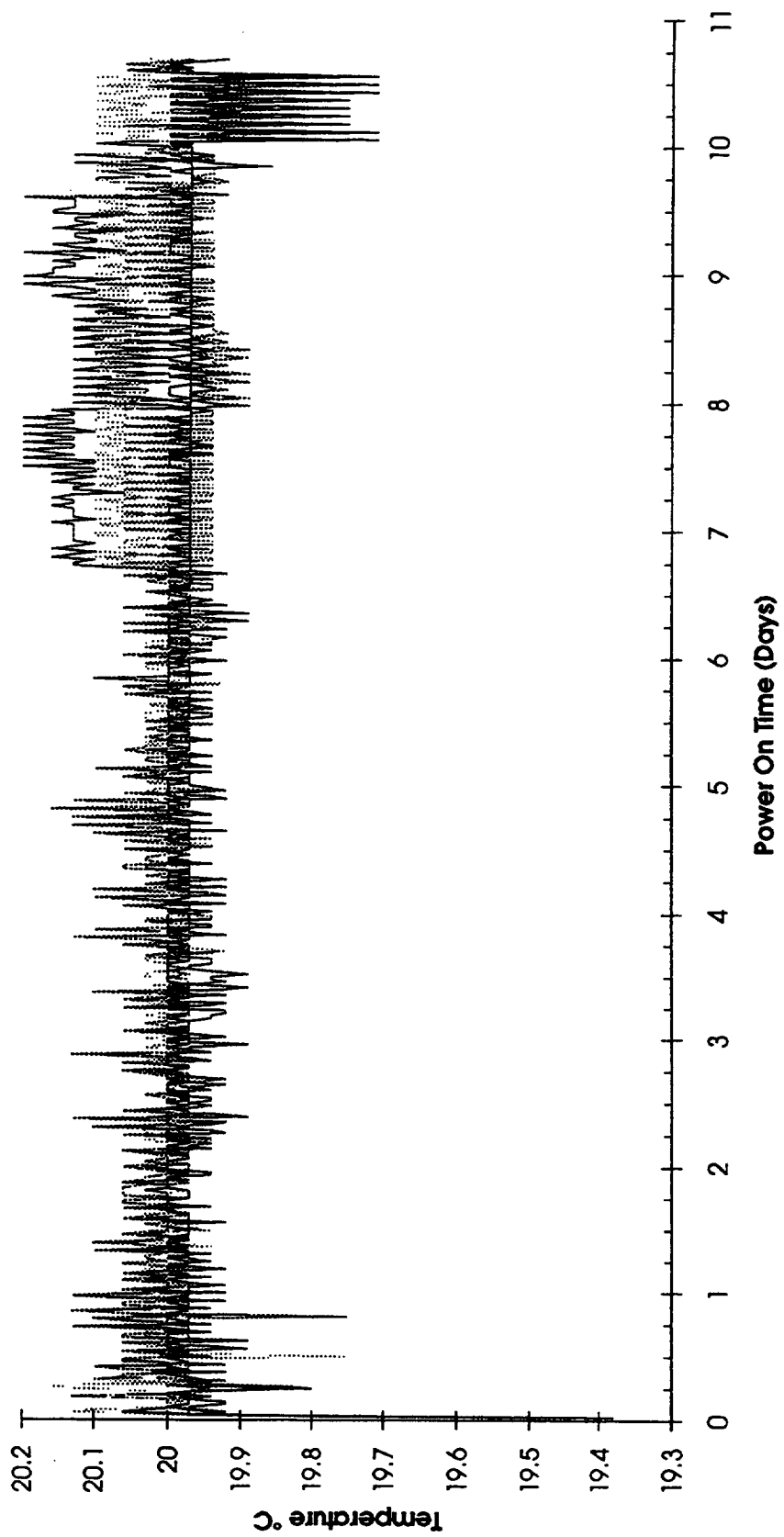


Figure 12. Chamber Bottom Thermistor Temperatures (Fine Channels 7-10)

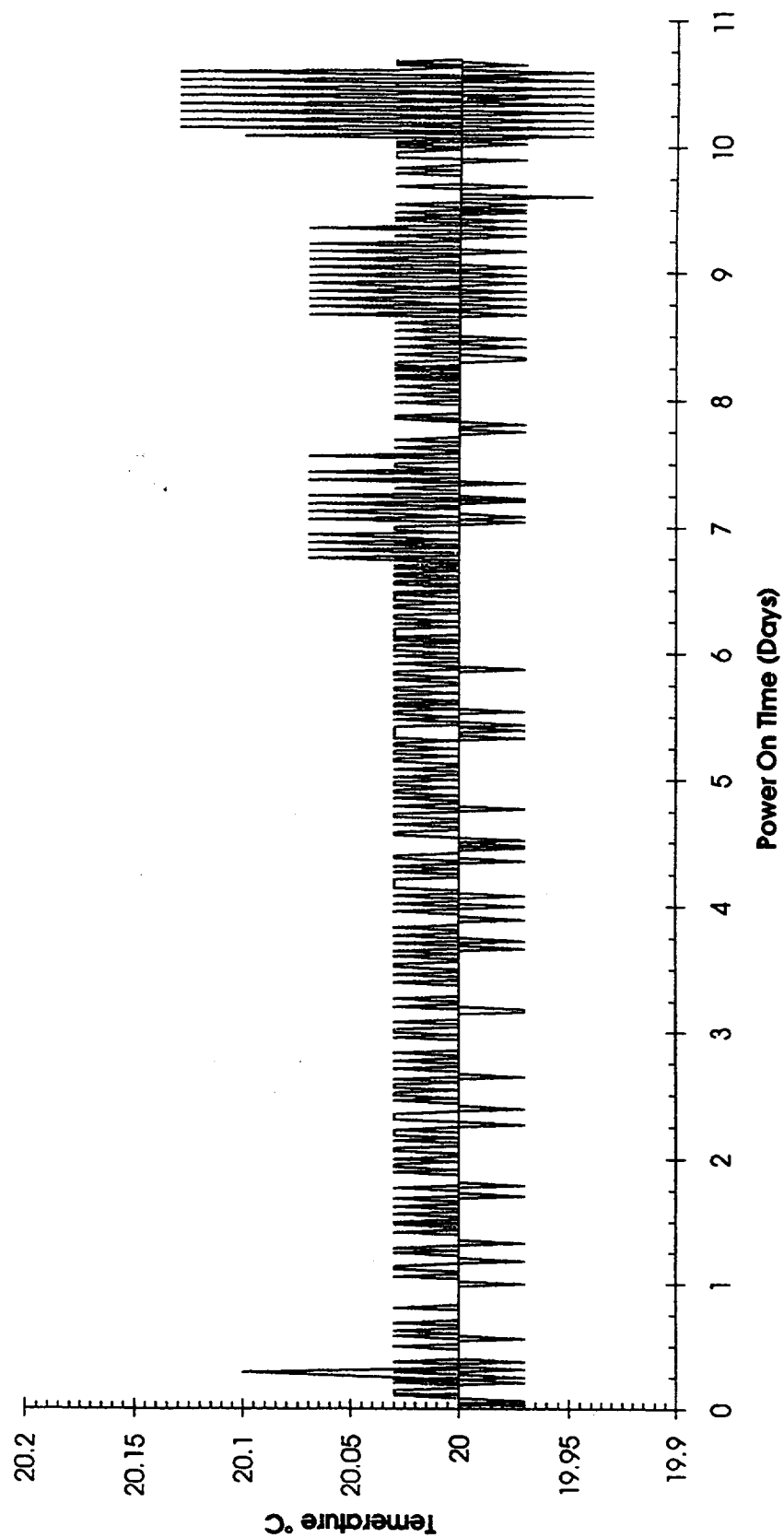


Figure 13. Electronics Unit Thermistor Temperature (Fine Channel 11)

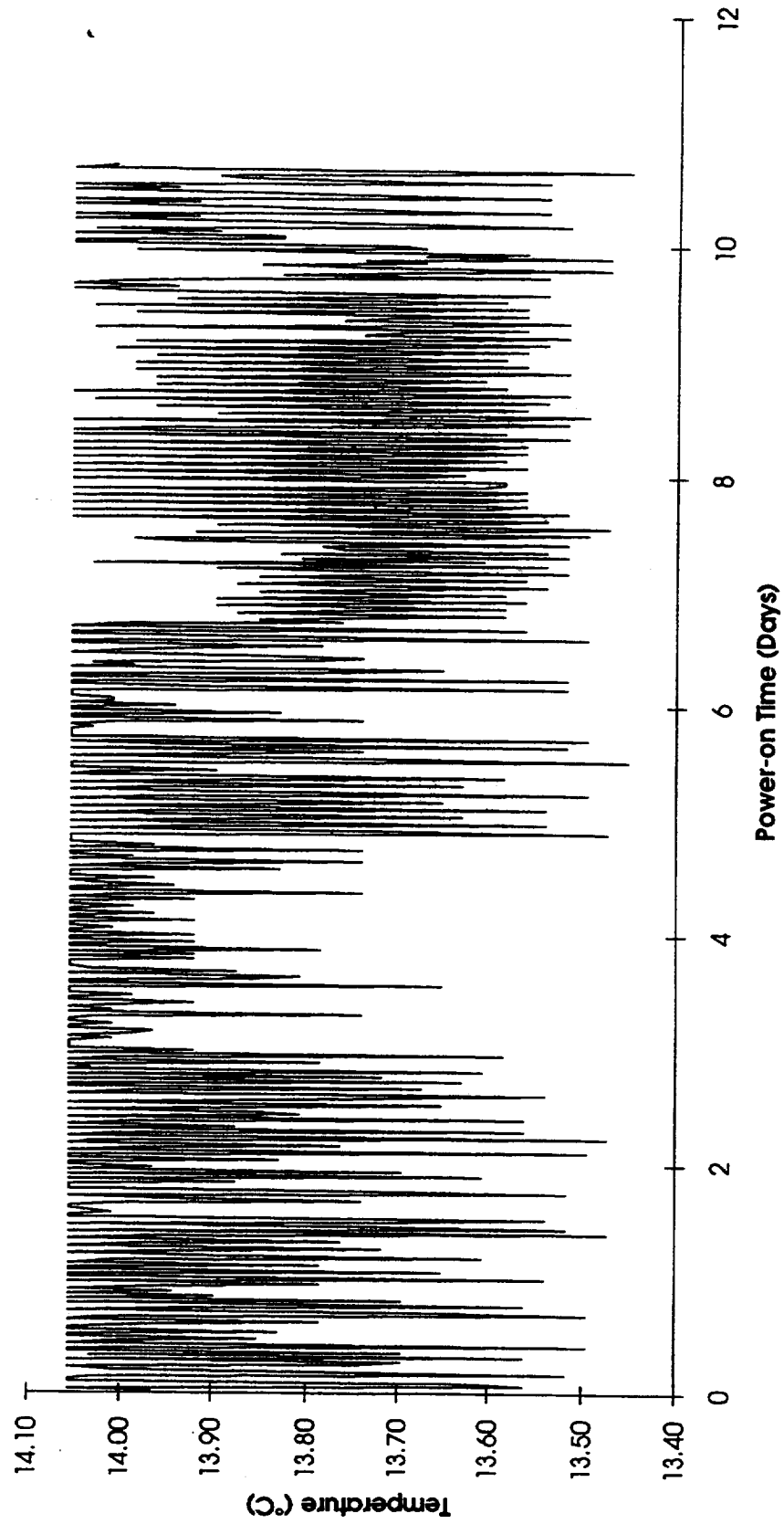


Figure 14. TempMentor Data for 3/3/94 to 3/19/94

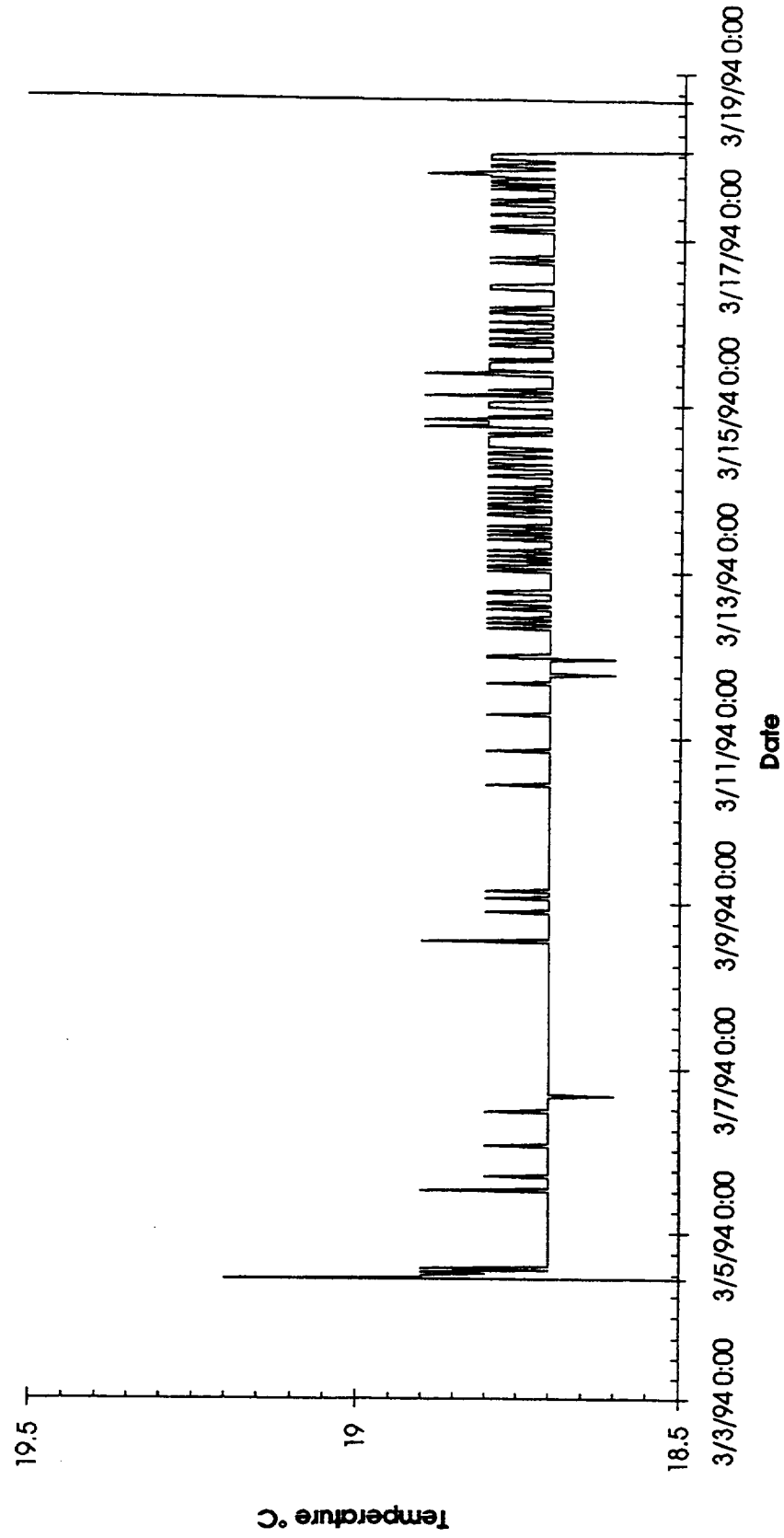
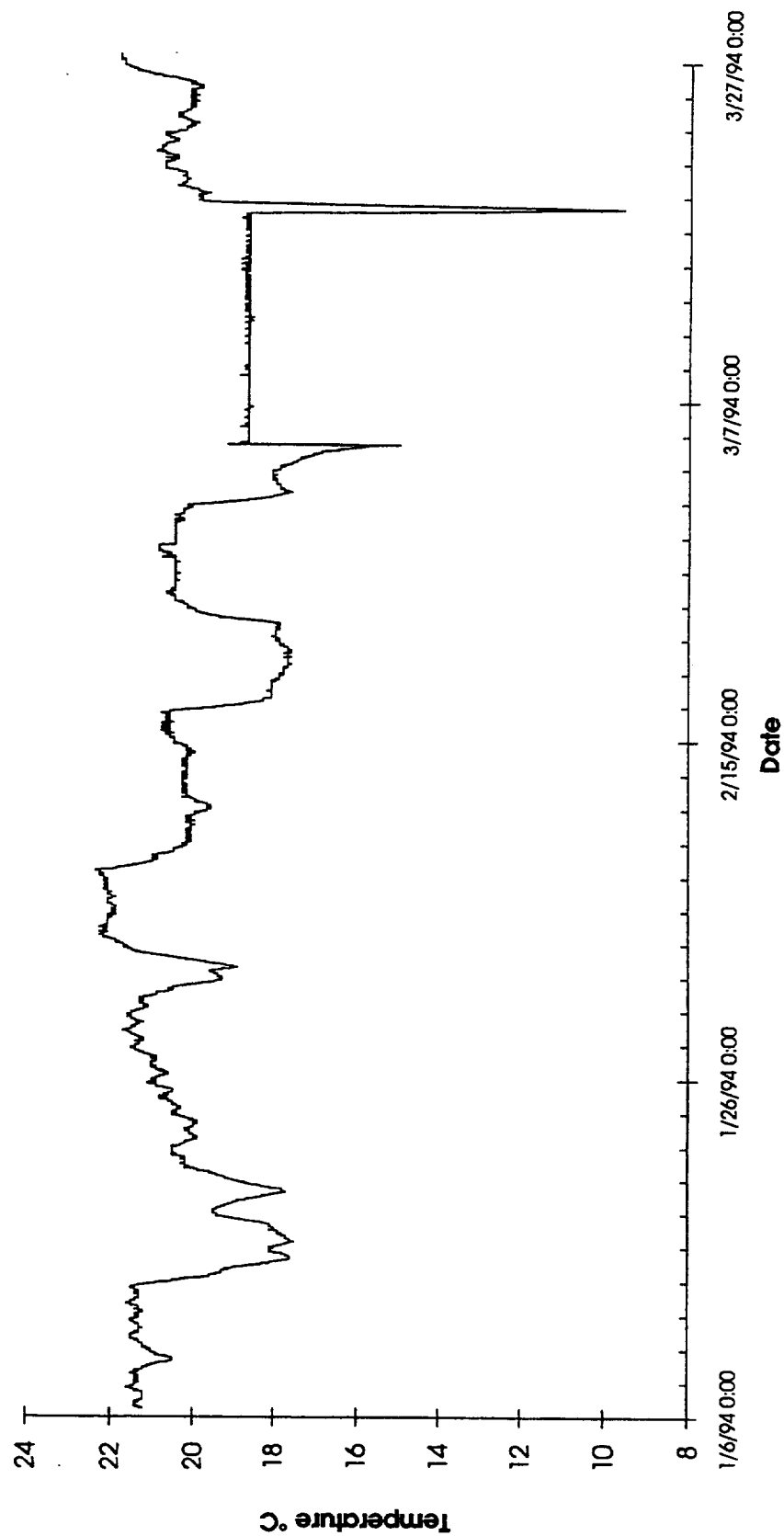


Figure 15. TempMentor Data for 1/6/94 to 3/27/94



4.3.2 Electronics Unit

The 7/93 version of the ECT flight software was used for the OAST-2/STS-62 mission. It was loaded into the pedigreed flight PROMs for the final assembly of the Electronics Unit. The flight software was verified by more than 150 hours of test operation before final assembly of the Electronics Unit. The 7/93 version of the software prevents data overwriting so that no stored data is lost after all available on-board memory is used. However, data storage was not optimized in the 7/93 version and all available on-board memory was used by Flight Day 11. (The STS-62 flight lasted fourteen days.) Nonetheless, nominal temperature measurement and heater power control continued without interruption and the recorded data was recovered. The Temp Mentor data (Figures 14 and 15) and the telemetry data (Appendices E - I) confirm that thermal control was uninterrupted.

Data storage can be and should be made more efficient for use in subsequent missions.

4.3.3 Ground Software

The LabView application was used by John Weber (FS) to create software to display and capture telemetry data on spacecraft pointing, experiment power and three temperatures in real time. Due to late changes in mission planning for telemetry by JSC spacecraft pointing data was not received. The power and temperature data parameters were successfully displayed on screen and captured on hard disk.

4.3.4 Data Products

Utilization of the telemetry data captured by the ECT ground system turned out to be rather difficult, requiring a great amount of manual file manipulation. Snapshots of the telemetry data were printed at the POCC every thirty minutes. Plots of this data were provided by GSFC's thermal contractor and are included in Appendices E - I.

Utilization of the stored data recovered from the Electronics Unit and the Temp Mentor was straightforward. This data is shown in Figures 10 - 15. Averaging of power values did require some post-processing of the Electronics Unit data.

The primary NASA data product for telemetry is essentially a copy of the entire mission telemetry stream, accompanied by a index. This contains all telemetry data from the Orbiter and from the Hitchhiker Avionics. The necessary post-processing task was not attempted. This data has been supplied on nine-track magnetic tape and on CD-ROM.

5.0 Issues for Future Flights

The next flight for ECT will likely be of much longer duration than the fourteen-day flight of OAST-2/STS-62. With minor modifications, the hardware and software can be made ready for such a mission. Five issues must be addressed: (1) limited carrier spacecraft electrical power for thermal control; (2) limited electrical power for Temp Mentor power and Electronics Unit memory back-up power from ECT on-board power sources; (3) limited Temp Mentor and Electronics Unit memory for data storage; (4) the availability and compatibility of telemetry data; and (5) the availability and compatibility of electrical connectors.

5.1 Limited Carrier Spacecraft Power

With regard to thermal control, limitations on available carrier spacecraft electrical power will necessitate modifications to the ECT thermal control design, such as covering a portion of the radiator surface with MLI or thermal louvers. A different temperature set point for the Emulsion Chamber may also be needed, with possible impacts on flight software and thermal switches. A lower set point will affect the length of power interruption which may be tolerated without damage to the emulsion stack.

5.2 Limited ECT On-board Power

Back-up power for Electronics Unit memory was provided by two AA alkaline cells. The Temp Mentor power was provided by four AA alkaline cells. The adequacy of these power sources must be evaluated against missions of longer duration. The present number and/or type of cells may be inadequate.

Once again, if a new Temp Mentor is to be carried aboard the Electronics Unit, then the power loop-back must be disabled on the Temp Mentor's RS-232 port. This is intended to prevent power depletion.

5.3 Limited Memory

Data storage in Electronics Unit memory should be optimized. This was known before the OAST-2/STS-62 flight, but Chris Watson (UAH) was not able to complete the task in time for delivery to KSC. With the efficient use of available Electronics Unit memory, the frequency of data storage and perhaps the number of parameters stored might be adjusted such that the available Electronics Unit memory is sufficient for a mission of longer duration. The Electronics Unit design allows adding memory to the system, but a memory upgrade will result in more expense and probably more testing. The appropriate sample rate for the Temp Mentor must also be selected, based on the Temp Mentor available memory and longer mission durations.

5.4 Telemetry Data

During the flight of OAST-2/STS-62, data was provided to ground personnel via telemetry for experiment power and three temperatures. However, this data was handled and formatted by the Hitchhiker Avionics system prior to transmission. To enable the use of telemetry data during another flight, similar arrangements must be provided by the carrier spacecraft and compatible hardware/software resources must be made ready for ground monitoring. Close coordination between the ECT team and the appropriate authorities will be vital to the availability of telemetry data during any subsequent mission.

5.5 Electrical Connectors

The lack of available and compatible electrical connectors may present significant difficulties, especially if schedules are short. It is recommended that at least a flight-qualified mating connector for the Electronics Unit power feedthrough connector, J1, be located and made available as soon as possible.

5.6 Mechanical Interfaces

ECT is designed to be compatible with GSFC's Small Top Pallet. This interface consists of a grid of 3/8"-24 inserts on 7.000 cm centers to accept primary structural fasteners and a staggered grid of #10-32 lockplates for tie-down fasteners. For the OAST-2 mission, GSFC supplied all mounting fasteners. ECT supplied thermal standoff hardware and special parts designed to transmit shear loads.

For a future mission, a new mounting plate might be developed, if no suitable existing structure is available. Only a portion of the 3/8"-24 hole pattern and a small number of #10-32 holes for tie-downs would be required.

6.0 MSFC Contacts

Rickey Clements, Quality Assurance	544-7394
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Dr. Tom Parnell, Project Scientist	544-7690
David Siersma, Payload Integration & Testing	544-1325

APPENDIX C

Emulsion Stack Assembly Procedure

EMULSION CHAMBER ASSEMBLY PROCEDURE

ECT FLIGHT ARTICLE

1/28/93

CAUTION: DARKROOM CONDITIONS REQUIRED!

1. Clean the internal surfaces of the major flight structural elements of the Emulsion Chamber with alcohol. These parts are: Upper Chamber (P/N 4005), Lower Chamber (P/N 4006), Top (P/N 4012) and Bottom Plate (P/N 4013).
2. Mask and paint the internal surfaces of the major flight structural elements of the Emulsion Chamber with Krylon, Clear, No. 1303. Steps 17 and 18 of this procedure may be performed at this point.
3. Install Lower Chamber (with O-ring) on Bottom Plate. Align the Lower Chamber with the Bottom Plate and install mechanical fasteners. Install vacuum hardware. Install Lucite shims in Lower Chamber at two adjoining wall locations.
4. Build up calorimeter module emulsion stack on top of Bottom Plate inside Lower Chamber. Install the internal thermistor(s) with Eccobond 285/11. Placement will be as directed by the PI. Cut grooves in shims and plates for thermistor leads, if required. Install feedthrough connectors.
5. Install remaining Lucite shims in Lower Chamber, between the calorimeter module emulsion stack and the Lower Chamber wall at the two remaining locations opposite those shims previously installed. Insert additional Lucite or stainless steel shims until the desired fit is obtained.
6. Install window frame and mylar sheet on Lower Chamber. Connect vacuum source to Lower Chamber vacuum port.
7. Evacuate Lower Chamber and check calorimeter module emulsion stack height and flatness. Remove window frame and mylar sheet and adjust calorimeter module emulsion stack as needed.
8. Repeat Steps 6 and 7 of this procedure as necessary to obtain desired stable calorimeter module emulsion stack height and flatness.
9. Install steel divider plate on Lower Chamber.
10. Install Upper Chamber (with O-ring) on Lower Chamber. Align the Upper Chamber with the Lower Chamber and install mechanical fasteners. Install vacuum hardware. Install Lucite shims in Lower Chamber at two adjoining wall locations.
11. Build up producer module emulsion stack on top of steel divider plate inside Upper Chamber. Install the internal thermistor(s) with Eccobond 285/11. Placement will be as directed by the PI. Cut grooves in shims and plates for thermistor leads, if required. Install feedthrough connectors.
12. Install remaining Lucite shims in Upper Chamber, between the calorimeter module emulsion stack and the Upper Chamber wall at the two remaining locations opposite those shims previously installed. Insert additional Lucite or stainless steel shims until the desired fit is obtained.
13. Install window frame and mylar sheet on Upper Chamber. Connect vacuum source to Upper Chamber vacuum port.

14. Evacuate Upper Chamber and check producer module emulsion stack height and flatness. Remove window frame and mylar sheet and adjust producer module emulsion stack as needed.
15. Repeat Steps 13 and 14 of this procedure as necessary to obtain desired stable producer module emulsion stack height and flatness.
16. Install Top (with O-ring) on Upper Chamber. Align the Top with the Upper Chamber and install mechanical fasteners.
17. Clean the external surfaces of the major flight structural elements of the Emulsion Chamber with alcohol. These parts are: Upper Chamber (P/N 4005), Lower Chamber (P/N 4006), Top (P/N 4012) and Bottom Plate (P/N 4013).
18. Install Emulsion Chamber external thermal and electrical hardware on the major flight structural elements of the Emulsion Chamber.

APPENDIX D

Flight Emulsion Stack Configuration

ECT MATERIALS COMPOSITION

NAME	MATERIAL	THICKNESS	MASS	HEIGHT	mfp	r.l.	Sum mfp	Sum r.l.	Sum
	NAME	µm	g/cm2	µm	%	%	%	%	g/cm2
*****	APPENDICES			200,000			0.000	0.000	0.000
RUBBER	SHIM	4,890.000	1.298	200,000	0.148	0.865	0.148	0.865	1.298
PLASTIC #1	NEUTRON MON	600.000	0.085	195,110	0.102	0.210	0.249	1.076	1.383
PLASTIC #2	NEUTRON MON	600.000	0.085	194,510	0.102	0.210	0.351	1.286	1.468
PLASTIC #3	NEUTRON MON	600.000	0.085	193,910	0.102	0.210	0.453	1.496	1.554
PLASTIC #4	THERMISTA	600.000	0.085	193,310	0.102	0.210	0.555	1.706	1.639
PLASTIC #5	NEUTRON MON	600.000	0.085	192,040	0.102	0.210	0.657	1.916	1.724
PLASTIC #6	NEUTRON MON	600.000	0.085	191,440	0.102	0.210	0.759	2.127	1.809
PLASTIC #7	TLD'S	600.000	0.085	190,840	0.102	0.210	0.861	2.337	1.894
*****	END OF APP.			190,240			0.861	2.337	1.894
				190,240			0.861	2.337	1.894
				190,240			0.861	2.337	1.894
*****	*****			190,240			0.861	2.337	1.894
PRIMARY	MODULE			190,240			0.861	2.337	1.894
*****	*****			190,240			0.861	2.337	1.894
				190,240			0.861	2.337	1.894
				190,240			0.861	2.337	1.894
HRH-HR8 #1	SCREEN FILM	3,500.000	0.081	190,240	0.920	5.466	1.781	7.803	1.976
P1 (E33-25)	CR-39	600.000	0.087	186,740	0.102	0.210	1.883	8.013	2.063
G01	GLASSINE	25.000	0.005	186,140	0.005	0.004	1.888	8.017	2.068
P02 (P1) —	* EMULSION 7B	200.000	0.077	186,115	0.058	0.702	1.946	8.719	2.145
	* LUCITE 300	300.000	0.035	185,915	0.051	0.105	1.997	8.824	2.180
	* EMULSION 6B	200.000	0.077	185,615	0.058	0.702	2.054	9.525	2.257
G03 —	GLASSINE	25.000	0.005	185,415	0.005	0.004	2.059	9.529	2.262
P03 —	* EMULSION 7B	75.000	0.029	185,390	0.022	0.263	2.081	9.792	2.291
	* LUCITE 500	800.000	0.059	185,315	0.136	0.280	2.217	10.073	2.350
	* EMULSION 6B	75.000	0.029	184,515	0.022	0.263	2.239	10.336	2.379
G04 —	GLASSINE	25.000	0.005	184,440	0.005	0.004	2.244	10.340	2.384
P04 —	* EMULSION 7B	200.000	0.772	184,415	0.058	0.702	2.301	11.041	3.156
	* LUCITE BASE	300.000	0.035	184,215	0.051	0.105	2.352	11.147	3.191
	* EMULSION 6B	200.000	0.772	183,915	0.058	0.702	2.410	11.848	3.963
G05 —	GLASSINE	25.000	0.005	183,715	0.005	0.004	2.415	11.852	3.968
P06 —	* EMULSION 7B	75.000	0.029	183,690	0.022	0.263	2.437	12.115	3.997
	* LUCITE BASE	800.000	0.059	183,615	0.136	0.280	2.573	12.396	4.056
	* EMULSION 7B	75.000	0.029	182,815	0.022	0.263	2.594	12.659	4.085
G07 —	GLASSINE	25.000	0.005	182,740	0.005	0.004	2.599	12.663	4.090
P07 —	* EMULSION 7B	200.000	0.772	182,715	0.058	0.702	2.657	13.364	4.862
	* LUCITE BASE	300.000	0.035	182,515	0.051	0.105	2.708	13.469	4.897
	* EMULSION 6B	200.000	0.772	182,215	0.058	0.702	2.765	14.171	5.669
G08 —	GLASSINE	25.000	0.005	182,015	0.022	0.263	2.787	14.434	5.674
P09 —	* EMULSION 7B	75.000	0.029	181,990	0.022	0.263	2.809	14.697	5.703
	* LUCITE BASE	800.000	0.059	181,915	0.136	0.280	2.945	14.978	5.762
	* EMULSION 7B	75.000	0.029	181,115	0.022	0.263	2.966	15.241	5.791
				181,040			2.966	15.241	5.791
*****	*****			181,040			2.966	15.241	5.791
PRODUCER	MODULE			181,040			2.966	15.241	5.791
*****	*****			181,040			2.966	15.241	5.791
				181,040			2.966	15.241	5.791
	*****			181,040			2.966	15.241	5.791
	PRODUCER			181,040			2.966	15.241	5.791
	1ST CYCLE			181,040			2.966	15.241	5.791
	*****			181,040			2.966	15.241	5.791
				181,040			2.966	15.241	5.791
G10 —	GLASSINE	25.000	0.005	181,040	0.022	0.263	2.988	15.504	5.796
P10 —	* EMULSION 7B	200.000	0.772	181,015	0.058	0.702	3.045	16.205	6.568

ECT MATERIALS COMPOSITION

NAME	MATERIAL	THICKNESS	MASS	HEIGHT	mfp	r.l.	Sum mfp	Sum r.l.	Sum
	NAME	µm	g/cm2	µm	%	%	%	%	g/cm2
	* LUCITE BASE	300.000	0.035	180,815	0.051	0.105	3.096	16.311	6.604
	* EMULSION 7B	200.000	0.772	180,515	0.058	0.702	3.154	17.012	7.376
G11 —	GLASSINE	25.000	0.005	180,315	0.022	0.263	3.176	17.275	7.381
P11 —	* EMULSION 7B	75.000	0.029	180,290	0.022	0.263	3.197	17.538	7.410
	* LUCITE BASE	800.000	0.059	180,215	0.136	0.280	3.333	17.819	7.469
	* EMULSION 7B	75.000	0.029	179,415	0.022	0.263	3.355	18.082	7.498
G12 —	GLASSINE	25.000	0.005	179,340	0.005	0.004	3.360	18.086	7.503
P12 —	* EMULSION 7B	75.000	0.029	179,315	0.022	0.263	3.381	18.349	7.532
	* LUCITE BASE	800.000	0.059	179,240	0.136	0.280	3.517	18.629	7.591
	* EMULSION 7B	75.000	0.029	178,440	0.022	0.263	3.539	18.892	7.620
G13 —	GLASSINE	25.000	0.005	178,365	0.005	0.004	3.544	18.896	7.625
L1	LEAD	500.000	0.568	178,340	0.292	8.910	3.836	27.806	8.192
G14 —	GLASSINE	25.000	0.005	177,840	0.022	0.263	3.858	28.069	8.197
P13 —	* EMULSION 7B	75.000	0.029	177,815	0.022	0.263	3.879	28.332	8.226
	* LUCITE BASE	800.000	0.059	177,740	0.136	0.280	4.015	28.613	8.285
	* EMULSION 7B	75.000	0.029	176,940	0.022	0.263	4.037	28.876	8.314
G15 —	GLASSINE	25.000	0.005	176,865	0.005	0.004	4.042	28.880	8.319
CR1 (P14)	CR-39	600.000	0.087	176,840	0.104	0.215	4.146	29.094	8.406
G16 —	GLASSINE	25.000	0.005	176,240	0.005	0.004	4.151	29.098	8.411
P15 —	* EMULSION 7B	75.000	0.029	176,215	0.022	0.263	4.173	29.361	8.440
	* LUCITE BASE	800.000	0.059	176,140	0.136	0.280	4.309	29.641	8.499
	* EMULSION 7B	75.000	0.029	175,340	0.022	0.263	4.330	29.905	8.528
				175,265			4.330	29.905	8.528
	*****			175,265			4.330	29.905	8.528
	PRODUCER			175,265			4.330	29.905	8.528
	2ND CYCLE			175,265			4.330	29.905	8.528
	*****			175,265			4.330	29.905	8.528
				175,265			4.330	29.905	8.528
G17 —	GLASSINE	25.000	0.005	175,265	0.022	0.263	4.352	30.168	8.533
P16 —	* EMULSION 7B	200.000	0.772	175,240	0.058	0.702	4.410	30.869	9.305
	* LUCITE BASE	300.000	0.035	175,040	0.051	0.105	4.461	30.974	9.341
	* EMULSION 6B	200.000	0.772	174,740	0.058	0.702	4.518	31.676	10.113
G18 —	GLASSINE	25.000	0.005	174,540	0.022	0.263	4.540	31.939	10.118
P17 —	* EMULSION 7B	75.000	0.029	174,515	0.022	0.263	4.561	32.202	10.147
	* LUCITE BASE	800.000	0.059	174,440	0.136	0.280	4.697	32.483	10.206
	* EMULSION 7B	75.000	0.029	173,640	0.022	0.263	4.719	32.746	10.235
G19 —	GLASSINE	25.000	0.005	173,565	0.005	0.004	4.724	32.750	10.240
P18 —	* EMULSION 7B	75.000	0.029	173,540	0.022	0.263	4.746	33.013	10.269
	* LUCITE BASE	800.000	0.059	173,465	0.136	0.280	4.882	33.293	10.328
	* EMULSION 7B	75.000	0.029	172,665	0.022	0.263	4.903	33.556	10.357
G20 —	GLASSINE	25.000	0.005	172,590	0.005	0.004	4.908	33.560	10.362
L2	LEAD	500.000	0.568	172,565	0.292	8.910	5.201	42.470	10.929
G21 —	GLASSINE	25.000	0.005	172,065	0.022	0.263	5.222	42.733	10.934
P19 —	* EMULSION 7B	75.000	0.029	172,040	0.022	0.263	5.244	42.996	10.963
	* LUCITE BASE	800.000	0.059	171,965	0.136	0.280	5.380	43.276	11.022
	* EMULSION 7B	75.000	0.029	171,165	0.022	0.263	5.401	43.540	11.051
G22 —	GLASSINE 25	25.000	0.005	171,090	0.005	0.004	5.406	43.543	11.056
CR2 (P20)	CR-39	600.000	0.087	171,065	0.104	0.215	5.510	43.758	11.143
G23 —	GLASSINE	25.000	0.005	170,465	0.005	0.004	5.516	43.762	11.148
P21 —	* EMULSION 7B	75.000	0.029	170,440	0.022	0.263	5.537	44.025	11.177
	* LUCITE BASE	800.000	0.059	170,365	0.136	0.280	5.673	44.305	11.236
	* EMULSION 7B	75.000	0.029	169,565	0.022	0.263	5.695	44.568	11.265
				169,490			5.695	44.568	11.265
	*****			169,490			5.695	44.568	11.265
	PRODUCER			169,490			5.695	44.568	11.265

ECT MATERIALS COMPOSITION

NAME	MATERIAL	THICKNESS	MASS	HEIGHT	mfp	r.l.	Sum mfp	Sum r.l.	Sum
	NAME	µm	g/cm2	µm	%	%	%	%	g/cm2
	3RD CYCLE			169,490			5.695	44.568	11.265
	*****			169,490			5.695	44.568	11.265
				169,490			5.695	44.568	11.265
G24 —	GLASSINE	25.000	0.005	169,490	0.005	0.004	5.700	44.572	11.270
P22 —	* EMULSION 7B	200.000	0.772	169,465	0.058	0.702	5.757	45.274	12.042
	* LUCITE BASE	300.000	0.035	169,265	0.051	0.105	5.808	45.379	12.078
	* EMULSION 7B	200.000	0.772	168,965	0.058	0.702	5.866	46.081	12.850
G25 —	GLASSINE	25.000	0.005	168,765	0.005	0.004	5.871	46.085	12.855
P23 —	* EMULSION 7B	75.000	0.029	168,740	0.022	0.263	5.893	46.348	12.884
	* LUCITE BASE	800.000	0.059	168,665	0.136	0.280	6.029	46.628	12.943
	* EMULSION 7B	75.000	0.029	167,865	0.022	0.263	6.050	46.891	12.972
G26 —	GLASSINE	25.000	0.005	167,790	0.005	0.004	6.055	46.895	12.977
P24 —	* EMULSION 7B	75.000	0.029	167,765	0.022	0.263	6.077	47.158	13.006
	* LUCITE BASE	800.000	0.059	167,690	0.136	0.280	6.213	47.439	13.065
	* EMULSION 7B	75.000	0.029	166,890	0.022	0.263	6.234	47.702	13.094
G27 —	GLASSINE	25.000	0.005	166,815	0.005	0.004	6.240	47.706	13.099
L3	LEAD	500.000	0.568	166,790	0.292	8.910	6.532	56.615	13.666
G28 —	GLASSINE	25.000	0.005	166,290	0.022	0.263	6.553	56.879	13.671
P25 —	* EMULSION 7B	75.000	0.029	166,265	0.022	0.263	6.575	57.142	13.700
	* LUCITE BASE	800.000	0.059	166,190	0.136	0.280	6.711	57.422	13.759
	* EMULSION 7B	75.000	0.029	165,390	0.022	0.263	6.733	57.685	13.788
G29 —	GLASSINE	25.000	0.005	165,315	0.005	0.004	6.738	57.689	13.793
CR3 (P26)	CR-39	600.000	0.087	165,290	0.104	0.215	6.842	57.904	13.880
G30 —	GLASSINE	25.000	0.005	164,690	0.005	0.004	6.847	57.908	13.885
P27 —	* EMULSION 7B	75.000	0.029	164,665	0.022	0.263	6.869	58.171	13.914
	* LUCITE BASE	800.000	0.059	164,590	0.136	0.280	7.004	58.451	13.973
	* EMULSION 7B	75.000	0.029	163,790	0.022	0.263	7.026	58.714	14.002
				163,715			7.026	58.714	14.002
	*****			163,715			7.026	58.714	14.002
	PRODUCER			163,715			7.026	58.714	14.002
	4TH CYCLE			163,715			7.026	58.714	14.002
	*****			163,715			7.026	58.714	14.002
				163,715			7.026	58.714	14.002
G31 —	GLASSINE	25.000	0.005	163,715	0.005	0.004	7.031	58.718	14.007
P28 —	* EMULSION 7B	200.000	0.772	163,690	0.058	0.702	7.089	59.420	14.779
	* LUCITE BASE	300.000	0.035	163,490	0.051	0.105	7.140	59.525	14.815
	* EMULSION 6B	200.000	0.772	163,190	0.058	0.702	7.197	60.226	15.587
G32 —	GLASSINE	25.000	0.005	162,990	0.005	0.004	7.202	60.230	15.592
P29 —	* EMULSION 7B	75.000	0.029	162,965	0.022	0.263	7.224	60.494	15.621
	* LUCITE BASE	800.000	0.059	162,890	0.136	0.280	7.360	60.774	15.680
	* EMULSION 7B	75.000	0.029	162,090	0.022	0.263	7.382	61.037	15.709
G33 —	GLASSINE	25.000	0.005	162,015	0.005	0.004	7.387	61.041	15.714
P30 —	* EMULSION 7B	75.000	0.029	161,990	0.022	0.263	7.408	61.304	15.743
	* LUCITE BASE	800.000	0.059	161,915	0.136	0.280	7.544	61.584	15.802
	* EMULSION 7B	75.000	0.029	161,115	0.022	0.263	7.566	61.847	15.831
G34 —	GLASSINE	25.000	0.005	161,040	0.005	0.004	7.571	61.851	15.836
L4	LEAD	500.000	0.568	161,015	0.292	8.910	7.863	70.761	16.403
G35 —	GLASSINE	25.000	0.005	160,515	0.022	0.263	7.885	71.024	16.408
P31 —	* EMULSION 7B	75.000	0.029	160,490	0.022	0.263	7.906	71.287	16.437
	* LUCITE BASE	800.000	0.059	160,415	0.136	0.280	8.042	71.568	16.496
	* EMULSION 7B	75.000	0.029	159,615	0.022	0.263	8.064	71.831	16.525
G36 —	GLASSINE	25.000	0.005	159,540	0.005	0.004	8.069	71.835	16.530
CR4 (P32)	CR-39	600.000	0.087	159,515	0.104	0.215	8.173	72.049	16.617
G37 —	GLASSINE	25.000	0.005	158,915	0.005	0.004	8.178	72.053	16.622
P33 —	* EMULSION 7B	75.000	0.029	158,890	0.022	0.263	8.200	72.316	16.651

ECT MATERIALS COMPOSITION

NAME	MATERIAL	THICKNESS	MASS	HEIGHT	mfp	r.l.	Sum mfp	Sum r.l.	Sum
	NAME	μm	g/cm2	μm	%	%	%	%	g/cm2
	* LUCITE BASE	800.000	0.059	158,815	0.136	0.280	8.336	72.597	16.710
	* EMULSION 7B	75.000	0.029	158,015	0.022	0.263	8.357	72.860	16.739
G37A	GLASSINE	25.000	0.005	157,940	0.005	0.004	8.362	72.864	16.744
E21-24	CR-39	600.000	0.087	157,915	0.104	0.215	8.467	73.078	16.831
G37B	GLASSINE	25.000	0.005	157,315	0.005	0.004	8.472	73.082	16.836
				157,290			8.472	73.082	16.836
	*****			157,290			8.472	73.082	16.836
	PRODUCER			157,290			8.472	73.082	16.836
	5TH CYCLE			157,290			8.472	73.082	16.836
	*****			157,290			8.472	73.082	16.836
				157,290			8.472	73.082	16.836
G38 —	GLASSINE	25.000	0.005	157,290	0.005	0.004	8.477	73.086	16.841
P34 —	* EMULSION 7B	200.000	0.772	157,265	0.058	0.702	8.534	73.788	17.613
	* LUCITE BASE	300.000	0.035	157,065	0.051	0.105	8.585	73.893	17.648
	* EMULSION 7B	200.000	0.772	156,765	0.058	0.702	8.643	74.595	18.420
G39 —	GLASSINE	25.000	0.005	156,565	0.005	0.004	8.648	74.598	18.425
P35 —	* EMULSION 7B	75.000	0.029	156,540	0.022	0.263	8.670	74.862	18.454
	* LUCITE BASE	800.000	0.059	156,465	0.136	0.280	8.806	75.142	18.513
	* EMULSION 7B	75.000	0.029	155,665	0.022	0.263	8.827	75.405	18.542
G40 —	GLASSINE	25.000	0.005	155,590	0.005	0.004	8.832	75.409	18.547
P36 —	* EMULSION 7B	75.000	0.029	155,565	0.022	0.263	8.854	75.672	18.576
	* LUCITE BASE	800.000	0.059	155,490	0.136	0.280	8.990	75.952	18.635
	* EMULSION 7B	75.000	0.029	154,690	0.022	0.263	9.012	76.215	18.664
G41 —	GLASSINE	25.000	0.005	154,615	0.005	0.004	9.017	76.219	18.669
L5	LEAD	500.000	0.568	154,590	0.292	8.910	9.309	85.129	19.237
G42 —	GLASSINE	25.000	0.005	154,090	0.022	0.263	9.330	85.392	19.242
P37 —	* EMULSION 7B	75.000	0.029	154,065	0.022	0.263	9.352	85.655	19.271
	* LUCITE BASE	800.000	0.059	153,990	0.136	0.280	9.488	85.936	19.330
	* EMULSION 7B	75.000	0.029	153,190	0.022	0.263	9.510	86.199	19.359
G43 —	GLASSINE	25.000	0.005	153,115	0.005	0.004	9.515	86.203	19.364
CR5 (P38)	CR-39	600.000	0.087	153,090	0.104	0.215	9.619	86.417	19.451
G44 —	GLASSINE	25.000	0.005	152,490	0.005	0.004	9.624	86.421	19.456
P39 —	* EMULSION 7B	75.000	0.029	152,465	0.022	0.263	9.646	86.684	19.485
	* LUCITE BASE	800.000	0.059	152,390	0.136	0.280	9.781	86.965	19.544
	* EMULSION 7B	75.000	0.029	151,590	0.022	0.263	9.803	87.228	19.573
				151,515			9.803	87.228	19.573
	*****			151,515			9.803	87.228	19.573
	PRODUCER			151,515			9.803	87.228	19.573
	6TH CYCLE			151,515			9.803	87.228	19.573
	*****			151,515			9.803	87.228	19.573
				151,515			9.803	87.228	19.573
G45 —	GLASSINE	25.000	0.005	151,515	0.005	0.004	9.808	87.232	19.578
P40 —	* EMULSION 7B	200.000	0.772	151,490	0.058	0.702	9.866	87.933	20.350
	* LUCITE BASE	300.000	0.035	151,290	0.051	0.105	9.917	88.038	20.385
	* EMULSION 6B	200.000	0.772	150,990	0.058	0.702	9.974	88.740	21.157
G46 —	GLASSINE	25.000	0.005	150,790	0.005	0.004	9.980	88.744	21.162
P41 —	* EMULSION 7B	75.000	0.029	150,765	0.022	0.263	10.001	89.007	21.191
	* LUCITE BASE	800.000	0.059	150,690	0.136	0.280	10.137	89.287	21.250
	* EMULSION 7B	75.000	0.029	149,890	0.022	0.263	10.159	89.551	21.279
G47 —	GLASSINE	25.000	0.005	149,815	0.005	0.004	10.164	89.554	21.284
P42 —	* EMULSION 7B	75.000	0.029	149,790	0.022	0.263	10.185	89.818	21.313
	* LUCITE BASE	800.000	0.059	149,715	0.136	0.280	10.321	90.098	21.372
	* EMULSION 7B	75.000	0.029	149,640	0.022	0.263	10.343	90.361	21.401
G48 —	GLASSINE	25.000	0.005	149,565	0.005	0.004	10.348	90.365	21.406
L6	LEAD	500.000	0.568	149,540	0.292	8.910	10.640	99.275	21.974

ECT MATERIALS COMPOSITION

NAME	MATERIAL	THICKNESS	MASS	HEIGHT	mfp	r.l.	Sum mfp	Sum r.l.	Sum
	NAME	µm	g/cm2	µm	%	%	%	%	g/cm2
G49 —	GLASSINE	25.000	0.005	149,040	0.022	0.263	10.662	99.538	21.979
P43 —	* EMULSION 7B	75.000	0.029	149,015	0.022	0.263	10.683	99.801	22.008
	* LUCITE BASE	800.000	0.059	148,940	0.136	0.280	10.819	100.081	22.067
	* EMULSION 7B	75.000	0.029	148,140	0.022	0.263	10.841	100.344	22.096
G50 —	GLASSINE	25.000	0.005	148,065	0.005	0.004	10.846	100.348	22.101
CR6(P50)	CR-39	600.000	0.087	148,040	0.104	0.215	10.950	100.563	22.188
G51 —	GLASSINE	25.000	0.005	147,440	0.005	0.004	10.955	100.567	22.193
P45 —	* EMULSION 7B	75.000	0.029	147,415	0.022	0.263	10.977	100.830	22.222
	* LUCITE BASE	800.000	0.059	147,340	0.136	0.280	11.113	101.110	22.281
	* EMULSION 7B	75.000	0.029	146,540	0.022	0.263	11.134	101.373	22.310
				146,465			11.134	101.373	22.310
	*****			146,465			11.134	101.373	22.310
	PRODUCER			146,465			11.134	101.373	22.310
	7TH CYCLE			146,465			11.134	101.373	22.310
	*****			146,465			11.134	101.373	22.310
				146,465			11.134	101.373	22.310
G52 —	GLASSINE	25.000	0.005	146,465	0.005	0.004	11.140	101.377	22.315
P46 —	* EMULSION 7B	200.000	0.772	146,440	0.058	0.702	11.197	102.079	23.087
	* LUCITE BASE	300.000	0.035	146,240	0.051	0.105	11.248	102.184	23.122
	* EMULSION 7B	200.000	0.772	145,940	0.058	0.702	11.306	102.886	23.894
G53 —	GLASSINE	25.000	0.005	145,740	0.005	0.004	11.311	102.890	23.899
P47 —	* EMULSION 7B	75.000	0.029	145,715	0.022	0.263	11.332	103.153	23.928
	* LUCITE BASE	800.000	0.059	145,640	0.136	0.280	11.468	103.433	23.987
	* EMULSION 7B	75.000	0.029	144,840	0.022	0.263	11.490	103.696	24.016
G54 —	GLASSINE	25.000	0.005	144,765	0.005	0.004	11.495	103.700	24.021
P48 —	* EMULSION 7B	75.000	0.029	144,740	0.022	0.263	11.517	103.963	24.050
	* LUCITE BASE	800.000	0.059	144,665	0.136	0.280	11.653	104.243	24.109
	* EMULSION 7B	75.000	0.029	143,865	0.022	0.263	11.674	104.507	24.138
G55 —	GLASSINE	25.000	0.005	143,790	0.005	0.004	11.679	104.510	24.143
L7	LEAD	500.000	0.568	143,765	0.292	8.910	11.972	113.420	24.711
G56 —	GLASSINE	25.000	0.005	143,265	0.022	0.263	11.993	113.683	24.716
P49 —	* EMULSION 7B	75.000	0.029	143,240	0.022	0.263	12.015	113.946	24.745
	* LUCITE BASE	800.000	0.059	143,165	0.136	0.280	12.151	114.227	24.804
	* EMULSION 7B	75.000	0.029	142,365	0.022	0.263	12.172	114.490	24.833
G57 —	GLASSINE	25.000	0.005	142,290	0.005	0.004	12.177	114.494	24.838
CR7(P50)E38	CR-39	600.000	0.087	142,265	0.104	0.215	12.281	114.708	24.925
G58 —	GLASSINE	25.000	0.005	141,665	0.005	0.004	12.287	114.712	24.930
P51 —	* EMULSION 7B	75.000	0.029	141,640	0.022	0.263	12.308	114.975	24.959
	* LUCITE BASE	800.000	0.059	141,565	0.136	0.280	12.444	115.256	25.018
	* EMULSION 7B	75.000	0.029	140,765	0.022	0.263	12.466	115.519	25.047
				140,690			12.466	115.519	25.047
	*****			140,690			12.466	115.519	25.047
	PRODUCER			140,690			12.466	115.519	25.047
	8TH CYCLE			140,690			12.466	115.519	25.047
	*****			140,690			12.466	115.519	25.047
				140,690			12.466	115.519	25.047
G59 —	GLASSINE	25.000	0.005	140,690	0.005	0.004	12.471	115.523	25.052
P52 —	* EMULSION 7B	200.000	0.772	140,665	0.058	0.702	12.528	116.224	25.824
	* LUCITE BASE	300.000	0.035	140,465	0.051	0.105	12.579	116.330	25.859
	* EMULSION 6B	200.000	0.772	140,165	0.058	0.702	12.637	117.031	26.631
G60 —	GLASSINE	25.000	0.005	139,965	0.005	0.004	12.642	117.035	26.636
P53 —	* EMULSION 7B	75.000	0.029	139,940	0.022	0.263	12.664	117.298	26.665
	* LUCITE BASE	800.000	0.059	139,865	0.136	0.280	12.800	117.579	26.724
	* EMULSION 7B	75.000	0.029	139,065	0.022	0.263	12.821	117.842	26.753
G61 —	GLASSINE	25.000	0.005	138,990	0.005	0.004	12.826	117.846	26.758

ECT MATERIALS COMPOSITION

NAME	MATERIAL	THICKNESS	MASS	HEIGHT	mfp	r.l.	Sum mfp	Sum r.l.	Sum
	NAME	µm	g/cm2	µm	%	%	%	%	g/cm2
P54	* EMULSION 7B	75.000	0.029	138,965	0.022	0.263	12.848	118.109	26.787
	* LUCITE BASE	800.000	0.059	138,890	0.136	0.280	12.984	118.389	26.846
	* EMULSION 7B	75.000	0.029	138,090	0.022	0.263	13.005	118.652	26.875
G62	GLASSINE	25.000	0.005	138,015	0.005	0.004	13.011	118.656	26.880
L8	LEAD	500.000	0.568	137,990	0.292	8.910	13.303	127.566	27.448
G63	GLASSINE	25.000	0.005	137,490	0.022	0.263	13.324	127.829	27.453
P55	* EMULSION 7B	75.000	0.029	137,465	0.022	0.263	13.346	128.092	27.482
	* LUCITE BASE	800.000	0.059	137,390	0.136	0.280	13.482	128.372	27.541
	* EMULSION 7B	75.000	0.029	136,590	0.022	0.263	13.504	128.635	27.570
G64	GLASSINE	25.000	0.005	136,515	0.005	0.004	13.509	128.639	27.575
CR8(P56)E30	CR-39	600.000	0.087	136,490	0.104	0.215	13.613	128.854	27.662
G65	GLASSINE	25.000	0.005	135,890	0.005	0.004	13.618	128.858	27.667
P57	* EMULSION 7B	75.000	0.029	135,865	0.022	0.263	13.640	129.121	27.696
	* LUCITE BASE	800.000	0.059	135,790	0.136	0.280	13.775	129.401	27.755
	* EMULSION 7B	75.000	0.029	134,990	0.022	0.263	13.797	129.664	27.784
				134,915			13.797	129.664	27.784
	*****			134,915			13.797	129.664	27.784
	PRODUCER			134,915			13.797	129.664	27.784
	9TH CYCLE			134,915			13.797	129.664	27.784
	*****			134,915			13.797	129.664	27.784
				134,915			13.797	129.664	27.784
G66	GLASSINE	25.000	0.005	134,915	0.005	0.004	13.802	129.668	27.789
P58	* EMULSION 7B	200.000	0.772	134,890	0.058	0.702	13.860	130.370	28.561
	* LUCITE BASE	300.000	0.035	134,690	0.051	0.105	13.911	130.475	28.596
	* EMULSION 7B	200.000	0.772	134,390	0.058	0.702	13.968	131.177	29.368
G67	GLASSINE	25.000	0.005	134,190	0.005	0.004	13.973	131.181	29.373
P59	* EMULSION 7B	75.000	0.029	134,165	0.022	0.263	13.995	131.444	29.402
	* LUCITE BASE	800.000	0.059	134,090	0.136	0.280	14.131	131.724	29.461
	* EMULSION 7B	75.000	0.029	133,290	0.022	0.263	14.153	131.987	29.490
G68	GLASSINE	25.000	0.005	133,240	0.005	0.004	14.158	131.991	29.495
P60	* EMULSION 7B	75.000	0.029	133,215	0.022	0.263	14.179	132.254	29.524
	* LUCITE BASE	800.000	0.059	133,140	0.136	0.280	14.315	132.535	29.583
	* EMULSION 7B	75.000	0.029	132,340	0.022	0.263	14.337	132.798	29.612
G69	GLASSINE	25.000	0.005	132,265	0.005	0.004	14.342	132.802	29.617
L9	LEAD	500.000	0.568	132,240	0.292	8.910	14.634	141.711	30.185
G70	GLASSINE	25.000	0.005	131,740	0.022	0.263	14.656	141.974	30.190
P61	* EMULSION 7B	75.000	0.029	131,715	0.022	0.263	14.677	142.238	30.219
	* LUCITE BASE	800.000	0.059	131,640	0.136	0.280	14.813	142.518	30.278
	* EMULSION 7B	75.000	0.029	130,840	0.022	0.263	14.835	142.781	30.307
G71	GLASSINE	25.000	0.005	130,765	0.005	0.004	14.840	142.785	30.312
CR9(P62)EC	CR-39	600.000	0.087	130,740	0.104	0.215	14.944	143.000	30.399
G72	GLASSINE	25.000	0.005	130,140	0.005	0.004	14.949	143.003	30.404
P63	* EMULSION 7B	75.000	0.029	130,115	0.022	0.263	14.971	143.267	30.433
	* LUCITE BASE	800.000	0.059	130,040	0.136	0.280	15.107	143.547	30.492
	* EMULSION 7B	75.000	0.029	129,240	0.022	0.263	15.128	143.810	30.521
				129,165			15.128	143.810	30.521
	*****			129,165			15.128	143.810	30.521
	PRODUCER			129,165			15.128	143.810	30.521
	10TH CYCLE			129,165			15.128	143.810	30.521
	*****			129,165			15.128	143.810	30.521
				129,165			15.128	143.810	30.521
G73	GLASSINE	25.000	0.005	129,165	0.005	0.004	15.133	143.814	30.526
P64	* EMULSION 7B	200.000	0.772	129,140	0.058	0.702	15.191	144.516	31.298
	* LUCITE BASE	300.000	0.035	128,940	0.051	0.105	15.242	144.621	31.333
	* EMULSION 6B	200.000	0.772	128,640	0.058	0.702	15.300	145.322	32.105

ECT MATERIALS COMPOSITION

NAME	MATERIAL	THICKNESS	MASS	HEIGHT	mfp	r.l.	Sum mfp	Sum r.l.	Sum
	NAME	µm	g/cm2	µm	%	%	%	%	g/cm2
G74 —	GLASSINE	25.000	0.005	128,440	0.005	0.004	15.305	145.326	32.110
P65 —	* EMULSION 7B	75.000	0.029	128,415	0.022	0.263	15.326	145.589	32.139
	* LUCITE BASE	800.000	0.059	128,340	0.136	0.280	15.462	145.870	32.198
	* EMULSION 7B	75.000	0.029	127,540	0.022	0.263	15.484	146.133	32.227
G75 —	GLASSINE	25.000	0.005	127,465	0.005	0.004	15.489	146.137	32.232
P66 —	* EMULSION 7B	75.000	0.029	127,440	0.022	0.263	15.511	146.400	32.261
	* LUCITE BASE	800.000	0.059	127,365	0.136	0.280	15.647	146.680	32.320
	* EMULSION 7B	75.000	0.029	126,565	0.022	0.263	15.668	146.943	32.349
G76 —	GLASSINE	25.000	0.005	126,490	0.005	0.004	15.673	146.947	32.354
L10	LEAD	500.000	0.568	126,465	0.292	8.910	15.966	155.857	32.921
G77 —	GLASSINE	25.000	0.005	125,965	0.022	0.263	15.987	156.120	32.926
P67 —	* EMULSION 7B	75.000	0.029	125,940	0.022	0.263	16.009	156.383	32.955
	* LUCITE BASE	800.000	0.059	125,865	0.136	0.280	16.145	156.663	33.014
	* EMULSION 7B	75.000	0.029	125,065	0.022	0.263	16.166	156.927	33.043
G78 —	GLASSINE	25.000	0.005	124,990	0.005	0.004	16.171	156.930	33.048
CR10(P68)E16	CR-39	600.000	0.087	124,965	0.104	0.215	16.275	157.145	33.135
G79 —	GLASSINE	25.000	0.005	124,365	0.005	0.004	16.281	157.149	33.140
P69 —	* EMULSION 7B	75.000	0.029	124,340	0.022	0.263	16.302	157.412	33.169
	* LUCITE BASE	800.000	0.059	124,265	0.136	0.280	16.438	157.692	33.228
	* EMULSION 7B	75.000	0.029	123,465	0.022	0.263	16.460	157.956	33.257
G79A —	GLASSINE	25.000	0.050	123,390	0.005	0.004	16.465	157.959	33.307
P70	HRH-HR8 #2	3,500.000	0.812	123,365	0.920	5.466	17.385	163.426	34.119
				119,865			17.385	163.426	34.119
*****	*****			119,865			17.385	163.426	34.119
END PRODUCER MODULE				119,865			17.385	163.426	34.119
*****	*****			119,865			17.385	163.426	34.119
				119,865			17.385	163.426	34.119
*****	*****			119,865			17.385	163.426	34.119
DIVIDER PLATE	STAINLESS	3,175.000	2.449	119,865	1.894	18.053	19.279	181.479	36.568
*****	*****			116,690			19.279	181.479	36.568
				116,690			19.279	181.479	36.568
*****	*****			116,690			19.279	181.479	36.568
CALORIMETER	MODULE (I)			116,690			19.279	181.479	36.568
*****	*****			116,690			19.279	181.479	36.568
				116,690			19.279	181.479	36.568
RUBBER	SHIM	2,000.000	0.528	116,690	0.600	3.520	19.879	184.999	37.096
PLASTIC	N-MONITOR #2	600.000	0.085	114,690	0.102	0.210	19.981	185.209	37.181
PLASTIC	N-MONITOR #1	600.000	0.085	114,090	0.102	0.210	20.083	185.419	37.266
PLASTIC	SHIM	800.000	0.211	113,490	0.240	1.408	20.322	186.828	37.477
G80 —	GLASSINE	25.000	0.005	112,690	0.005	0.004	20.327	186.832	37.482
C00 —	* EMULSION 7B	200.000	0.772	112,665	0.058	0.702	20.385	187.533	38.254
	* LUCITE BASE	300.000	0.035	112,465	0.051	0.105	20.436	187.638	38.290
	* EMULSION 7B	200.000	0.772	112,165	0.058	0.702	20.494	188.340	39.062
				111,965			20.494	188.340	39.062
**1ST CYCLE				111,965			20.494	188.340	39.062
				111,965			20.494	188.340	39.062
G81 —	GLASSINE	25.000	0.005	111,965	0.005	0.004	20.499	188.344	39.067
LC1	LEAD	1,000.000	1.135	111,940	0.585	17.820	21.083	206.163	40.202
G82 —	GLASSINE	25.000	0.005	110,940	0.005	0.004	21.088	206.167	40.207
				110,915			21.088	206.167	40.207
X1A	X-RAY FILM	220.000	0.043	110,915	0.032	0.392	21.121	206.559	40.250
X1B	X-RAY FILM	220.000	0.043	110,695	0.032	0.392	21.153	206.951	40.293
G83 —	GLASSINE	25.000	0.005	110,475	0.005	0.004	21.158	206.955	40.298
C01 —	* EMULSION 7B	75.000	0.029	110,450	0.022	0.263	21.179	207.218	40.327
	* LUCITE BASE	800.000	0.059	110,375	0.136	0.280	21.315	207.498	40.386

ECT MATERIALS COMPOSITION

NAME	MATERIAL	THICKNESS	MASS	HEIGHT	mfp	r.l.	Sum mfp	Sum r.l.	Sum
	NAME	μm	g/cm2	μm	%	%	%	%	g/cm2
	* EMULSION 7B	75.000	0.029	109,575	0.022	0.263	21.337	207.761	40.415
				109,500			21.337	207.761	40.415
**2ND CYCLE				109,500			21.337	207.761	40.415
				109,500			21.337	207.761	40.415
G84 —	GLASSINE	25.000	0.005	109,500	0.005	0.004	21.342	207.765	40.420
LC2	LEAD	1,000.000	1.135	109,475	0.585	17.820	21.926	225.584	41.555
G85 —	GLASSINE	25.000	0.005	108,475	0.005	0.004	21.932	225.588	41.560
				108,450			21.932	225.588	41.560
X2A	X-RAY FILM	220.000	0.043	108,450	0.032	0.392	21.964	225.980	41.603
X2B	X-RAY FILM	220.000	0.043	108,230	0.032	0.392	21.996	226.372	41.646
G86 —	GLASSINE	25.000	0.005	108,010	0.005	0.004	22.001	226.376	41.651
C02 —	* EMULSION 7B	75.000	0.029	107,985	0.022	0.263	22.023	226.639	41.680
	* LUCITE BASE	800.000	0.059	107,910	0.136	0.280	22.158	226.919	41.739
	* EMULSION 7B	75.000	0.029	107,110	0.022	0.263	22.180	227.182	41.768
				107,035			22.180	227.182	41.768
**3RD CYCLE				107,035			22.180	227.182	41.768
				107,035			22.180	227.182	41.768
G87 —	GLASSINE	25.000	0.005	107,035	0.005	0.004	22.185	227.186	41.773
LC3	LEAD	1,000.000	1.135	107,010	0.585	17.820	22.770	245.005	42.908
G88 —	GLASSINE	25.000	0.005	106,010	0.005	0.004	22.775	245.009	42.913
				105,985			22.775	245.009	42.913
X3A	X-RAY FILM	220.000	0.043	105,985	0.032	0.392	22.807	245.401	42.956
X3B	X-RAY FILM	220.000	0.043	105,765	0.032	0.392	22.839	245.793	42.999
G89 —	GLASSINE	25.000	0.005	105,545	0.005	0.004	22.844	245.797	43.004
C03 —	* EMULSION 7B	75.000	0.029	105,520	0.022	0.263	22.866	246.060	43.033
	* LUCITE BASE	800.000	0.059	105,445	0.136	0.280	23.002	246.340	43.092
	* EMULSION 7B	75.000	0.029	104,645	0.022	0.263	23.023	246.603	43.121
				104,570			23.023	246.603	43.121
**4TH CYCLE				104,570			23.023	246.603	43.121
				104,570			23.023	246.603	43.121
G90 —	GLASSINE	25.000	0.005	104,570	0.005	0.004	23.028	246.607	43.126
LC4	LEAD	1,000.000	1.135	104,545	0.585	17.820	23.613	264.426	44.261
G91 —	GLASSINE	25.000	0.005	103,545	0.005	0.004	23.618	264.430	44.266
				103,520			23.618	264.430	44.266
X4A	X-RAY FILM	220.000	0.043	103,520	0.032	0.392	23.650	264.822	44.309
X4B	X-RAY FILM	220.000	0.043	103,300	0.032	0.392	23.682	265.214	44.352
G92 —	GLASSINE	25.000	0.005	103,080	0.005	0.004	23.687	265.218	44.357
C04 —	* EMULSION 7B	75.000	0.029	103,055	0.022	0.263	23.709	265.481	44.386
	* LUCITE BASE	800.000	0.059	102,980	0.136	0.280	23.845	265.761	44.445
	* EMULSION 7B	75.000	0.029	102,180	0.022	0.263	23.866	266.024	44.474
				102,105			23.866	266.024	44.474
**5TH CYCLE				102,105			23.866	266.024	44.474
				102,105			23.866	266.024	44.474
G93 —	GLASSINE	25.000	0.005	102,105	0.005	0.004	23.872	266.028	44.479
LC5	LEAD	1,000.000	1.135	102,080	0.585	17.820	24.456	283.847	45.614
G94 —	GLASSINE	25.000	0.005	101,080	0.005	0.004	24.461	283.851	45.619
				101,055			24.461	283.851	45.619
X5A	X-RAY FILM	220.000	0.043	101,055	0.032	0.392	24.493	284.243	45.663
X5B	X-RAY FILM	220.000	0.043	100,835	0.032	0.392	24.525	284.635	45.706
G95 —	GLASSINE	25.000	0.005	100,615	0.005	0.004	24.531	284.639	45.711
C05 —	* EMULSION 7B	75.000	0.029	100,590	0.022	0.263	24.552	284.902	45.740
	* LUCITE BASE	800.000	0.059	100,515	0.136	0.280	24.688	285.182	45.799
	* EMULSION 7B	75.000	0.029	99,715	0.022	0.263	24.710	285.445	45.828
G96 —	GLASSINE	25.000	0.005	99,640	0.005	0.004	24.715	285.449	45.833
B3(E15-E14)	CR39	600.000	0.087	99,615	0.104	0.215	24.819	285.664	45.920

ECT MATERIALS COMPOSITION

NAME	MATERIAL	THICKNESS	MASS	HEIGHT	mfp	r.l.	Sum mfp	Sum r.l.	Sum
	NAME	µm	g/cm2	µm	%	%	%	%	g/cm2
				99,015			24.819	285.664	45.920
				99,015			24.819	285.664	45.920
*****	*****			99,015			24.819	285.664	45.920
CALORIMETER	MODULE (II)			99,015			24.819	285.664	45.920
*****	*****			99,015			24.819	285.664	45.920
(6TH - 8TH CYCLES OMITTED)				99,015			24.819	285.664	45.920
				99,015			24.819	285.664	45.920
**9TH CYCLE				99,015			24.819	285.664	45.920
				99,015			24.819	285.664	45.920
G97 —	GLASSINE	25.000	0.005	99,015	0.005	0.004	24.824	285.668	45.925
LC9	LEAD	2,000.000	2.271	98,990	1.170	35.655	25.994	321.322	48.196
G98 —	GLASSINE	25.000	0.005	96,990	0.005	0.004	25.999	321.326	48.201
				96,965			25.999	321.326	48.201
X9A	X-RAY FILM	220.000	0.043	96,965	0.032	0.392	26.031	321.718	48.244
X9B	X-RAY FILM	220.000	0.043	96,745	0.032	0.392	26.063	322.109	48.287
G99 —	GLASSINE	25.000	0.005	96,525	0.005	0.004	26.068	322.113	48.292
C09 —	* EMULSION 7B	75.000	0.029	96,500	0.022	0.263	26.090	322.376	48.321
	* LUCITE BASE	800.000	0.059	96,425	0.136	0.280	26.226	322.657	48.380
	* EMULSION 7B	75.000	0.029	95,625	0.022	0.263	26.247	322.920	48.409
G100 —	GLASSINE	25.000	0.005	95,550	0.005	0.004	26.252	322.924	48.414
B3(E15-E14)	CR39	600.000	0.087	94,950	0.104	0.215	26.356	323.138	48.501
				94,350			26.356	323.138	48.501
*10TH CYCLE				94,350			26.356	323.138	48.501
				94,350			26.356	323.138	48.501
G101 —	GLASSINE	25.000	0.005	94,350	0.005	0.004	26.361	323.142	48.506
LC10	LEAD	2,000.000	2.271	94,325	1.170	35.655	27.531	358.797	50.777
G102 —	GLASSINE	25.000	0.005	92,325	0.005	0.004	27.536	358.801	50.782
				92,300			27.536	358.801	50.782
X10A	X-RAY FILM	220.000	0.043	92,300	0.032	0.392	27.568	359.193	50.825
X10B	X-RAY FILM	220.000	0.043	92,080	0.032	0.392	27.600	359.584	50.868
G103 —	GLASSINE	25.000	0.005	91,860	0.005	0.004	27.606	359.588	50.873
C10 —	* EMULSION 7B	75.000	0.029	91,835	0.022	0.263	27.627	359.851	50.902
	* LUCITE BASE	800.000	0.059	91,760	0.136	0.280	27.763	360.131	50.961
	* EMULSION 7B	75.000	0.029	91,035	0.022	0.263	27.785	360.395	50.990
				90,960			27.785	360.395	50.990
*11TH CYCLE				90,960			27.785	360.395	50.990
				90,960			27.785	360.395	50.990
G104 —	GLASSINE	25.000	0.005	90,960	0.005	0.004	27.790	360.398	50.995
LC11	LEAD	2,000.000	2.271	90,935	1.170	35.655	28.959	396.053	53.266
G105 —	GLASSINE	25.000	0.005	88,935	0.005	0.004	28.965	396.057	53.271
				88,910			28.965	396.057	53.271
X11A	X-RAY FILM	220.000	0.043	88,910	0.032	0.392	28.997	396.449	53.314
X11B	X-RAY FILM	220.000	0.043	88,690	0.032	0.392	29.029	396.840	53.357
G106 —	GLASSINE	25.000	0.005	88,470	0.005	0.004	29.034	396.844	53.362
C11 —	* EMULSION 7B	75.000	0.029	88,445	0.022	0.263	29.055	397.107	53.391
	* LUCITE BASE	800.000	0.059	88,370	0.136	0.280	29.191	397.388	53.450
	* EMULSION 7B	75.000	0.029	87,570	0.022	0.263	29.213	397.651	53.479
				87,495			29.213	397.651	53.479
*12TH CYCLE				87,495			29.213	397.651	53.479
				87,495			29.213	397.651	53.479
G107 —	GLASSINE	25.000	0.005	87,495	0.005	0.004	29.218	397.655	53.484
LC12	LEAD	2,000.000	2.271	87,470	1.170	35.655	30.388	433.309	55.755
G108 —	GLASSINE	25.000	0.005	85,470	0.005	0.004	30.393	433.313	55.760
				85,445			30.393	433.313	55.760
X12A	X-RAY FILM	220.000	0.043	85,665	0.032	0.392	30.425	433.705	55.803

ECT MATERIALS COMPOSITION

NAME	MATERIAL	THICKNESS	MASS	HEIGHT	mfp	r.l.	Sum mfp	Sum r.l.	Sum
	NAME	µm	g/cm2	µm	%	%	%	%	g/cm2
X12B	X-RAY FILM	220.000	0.043	85,445	0.032	0.392	30.457	434.097	55.846
G109	GLASSINE	25.000	0.005	85,225	0.005	0.004	30.462	434.100	55.851
C12	* EMULSION 7B	75.000	0.029	85,200	0.022	0.263	30.484	434.364	55.880
	* LUCITE BASE	800.000	0.059	85,125	0.136	0.280	30.620	434.644	55.939
	* EMULSION 7B	75.000	0.029	84,325	0.022	0.263	30.641	434.907	55.968
				84,250			30.641	434.907	55.968
*13TH CYCLE				84,250			30.641	434.907	55.968
				84,250			30.641	434.907	55.968
G110	GLASSINE	25.000	0.005	84,250	0.005	0.004	30.646	434.911	55.973
LC13	LEAD	2,000.000	2.271	84,225	1.170	35.655	31.816	470.566	58.244
G111	GLASSINE	25.000	0.005	82,225	0.005	0.004	31.821	470.570	58.249
				82,200			31.821	470.570	58.249
X13A	X-RAY FILM	220.000	0.043	82,200	0.032	0.392	31.853	470.961	58.292
X13B	X-RAY FILM	220.000	0.043	81,980	0.032	0.392	31.885	471.353	58.335
G112	GLASSINE	25.000	0.005	81,760	0.005	0.004	31.890	471.357	58.340
C13	* EMULSION 7B	75.000	0.029	81,735	0.022	0.263	31.912	471.620	58.369
	* LUCITE BASE	800.000	0.059	81,660	0.136	0.280	32.048	471.900	58.428
	* EMULSION 7B	75.000	0.029	80,935	0.022	0.263	32.070	472.163	58.457
				80,860			32.070	472.163	58.457
*14TH CYCLE				80,860			32.070	472.163	58.457
				80,860			32.070	472.163	58.457
G113	GLASSINE	25.000	0.005	80,860	0.005	0.004	32.075	472.167	58.462
LC14	LEAD	2,000.000	2.271	80,835	1.170	35.655	33.244	507.822	60.733
G114	GLASSINE	25.000	0.005	78,835	0.005	0.004	33.249	507.826	60.738
				78,810			33.249	507.826	60.738
X14A	X-RAY FILM	220.000	0.043	78,810	0.032	0.392	33.281	508.217	60.781
X14B	X-RAY FILM	220.000	0.043	78,590	0.032	0.392	33.314	508.609	60.825
G115	GLASSINE	25.000	0.005	78,370	0.005	0.004	33.319	508.613	60.830
C14	* EMULSION 7B	75.000	0.029	78,345	0.022	0.263	33.340	508.876	60.859
	* LUCITE BASE	800.000	0.059	78,270	0.136	0.280	33.476	509.156	60.918
	* EMULSION 7B	75.000	0.029	77,470	0.022	0.263	33.498	509.419	60.947
G116	GLASSINE	25.000	0.005	77,395	0.005	0.004	33.503	509.423	60.952
B4(E19-E20)	CR-39	600.000	0.087	77,370	0.104	0.215	33.607	509.638	61.039
				76,770			33.607	509.638	61.039
*15TH CYCLE				76,770			33.607	509.638	61.039
				76,770			33.607	509.638	61.039
G117	GLASSINE	25.000	0.005	76,770	0.005	0.004	33.612	509.642	61.044
LC15	LEAD	2,000.000	2.271	76,745	1.170	35.655	34.782	545.297	63.315
G118	GLASSINE	25.000	0.005	74,745	0.005	0.004	34.787	545.301	63.320
				74,720			34.787	545.301	63.320
X15A	X-RAY FILM	220.000	0.043	74,720	0.032	0.392	34.819	545.692	63.363
X15B	X-RAY FILM	220.000	0.043	74,500	0.032	0.392	34.851	546.084	63.406
G119	GLASSINE	25.000	0.005	74,280	0.005	0.004	34.856	546.088	63.411
C15	* EMULSION 7B	75.000	0.029	74,255	0.022	0.263	34.878	546.351	63.440
	* LUCITE BASE	800.000	0.059	74,180	0.136	0.280	35.014	546.631	63.499
	* EMULSION 7B	75.000	0.029	73,380	0.022	0.263	35.035	546.894	63.528
				73,305			35.035	546.894	63.528
*16TH CYCLE				73,305			35.035	546.894	63.528
				73,305			35.035	546.894	63.528
G120	GLASSINE	25.000	0.005	73,280	0.005	0.004	35.040	546.898	63.533
LC16	LEAD + TLD	2,000.000	2.271	73,255	1.170	35.655	36.210	582.553	65.804
G121	GLASSINE	25.000	0.005	71,255	0.005	0.004	36.215	582.557	65.809
				71,230			36.215	582.557	65.809
X16A	X-RAY FILM	220.000	0.043	71,230	0.032	0.392	36.247	582.948	65.852
X16B	X-RAY FILM	220.000	0.043	71,010	0.032	0.392	36.279	583.340	65.895

ECT MATERIALS COMPOSITION

NAME	MATERIAL	THICKNESS	MASS	HEIGHT	mfp	r.l.	Sum mfp	Sum r.l.	Sum
	NAME	µm	g/cm2	µm	%	%	%	%	g/cm2
G122 --	GLASSINE	25.000	0.005	70,790	0.005	0.004	36.285	583.344	65.900
C16 -----	EMULSION 7B	75.000	0.029	70,765	0.022	0.263	36.306	583.607	65.929
	* LUCITE BASE	800.000	0.059	70,690	0.136	0.280	36.442	583.887	65.988
	* EMULSION 7B	75.000	0.029	69,890	0.022	0.263	36.464	584.150	66.017
				69,815			36.464	584.150	66.017
*17TH CYCLE				69,815			36.464	584.150	66.017
				69,815			36.464	584.150	66.017
G123 --	GLASSINE	25.000	0.005	69,815	0.005	0.004	36.469	584.154	66.022
LC17	LEAD	2,000.000	2.271	69,790	1.170	35.655	37.638	619.809	68.293
G124 --	GLASSINE	25.000	0.005	67,790	0.005	0.004	37.643	619.813	68.298
				67,765			37.643	619.813	68.298
X17A	X-RAY FILM	220.000	0.043	67,765	0.032	0.392	37.676	620.205	68.341
X17B	X-RAY FILM	220.000	0.043	67,545	0.032	0.392	37.708	620.596	68.384
G125 --	GLASSINE	25.000	0.005	67,325	0.005	0.004	37.713	620.600	68.389
C17 -----	EMULSION 7B	75.000	0.029	67,300	0.022	0.263	37.734	620.863	68.418
	* LUCITE BASE	800.000	0.059	67,225	0.136	0.280	37.870	621.143	68.477
	* EMULSION 7B	75.000	0.029	66,425	0.022	0.263	37.892	621.407	68.506
				66,350			37.892	621.407	68.506
*18TH CYCLE				66,350			37.892	621.407	68.506
				66,350			37.892	621.407	68.506
G126 --	GLASSINE	25.000	0.005	66,350	0.005	0.004	37.897	621.411	68.511
LC18	LEAD	2,000.000	2.271	66,325	1.170	35.655	39.067	657.065	70.782
G127 --	GLASSINE	25.000	0.005	64,325	0.005	0.004	39.072	657.069	70.787
				64,300			39.072	657.069	70.787
X18A	X-RAY FILM	220.000	0.043	64,300	0.032	0.392	39.104	657.461	70.830
X18B	X-RAY FILM	220.000	0.043	64,080	0.032	0.392	39.136	657.852	70.873
G128 --	GLASSINE	25.000	0.005	63,860	0.005	0.004	39.141	657.856	70.878
C18 -----	EMULSION 7B	75.000	0.029	63,835	0.022	0.263	39.163	658.119	70.907
	* LUCITE BASE	800.000	0.059	63,760	0.136	0.280	39.299	658.400	70.966
	* EMULSION 7B	75.000	0.029	62,960	0.022	0.263	39.320	658.663	70.995
				62,885			39.320	658.663	70.995
*19TH CYCLE				62,885			39.320	658.663	70.995
				62,885			39.320	658.663	70.995
G129 --	GLASSINE	25.000	0.005	62,885	0.005	0.004	39.325	658.667	71.000
LC19	LEAD	2,000.000	2.271	62,860	1.170	35.655	40.495	694.321	73.271
G130 --	GLASSINE	25.000	0.005	60,860	0.005	0.004	40.500	694.325	73.276
				60,835			40.500	694.325	73.276
X19A	X-RAY FILM	220.000	0.043	60,835	0.032	0.392	40.532	694.717	73.319
X19B	X-RAY FILM	220.000	0.043	60,615	0.032	0.392	40.564	695.109	73.362
G131 --	GLASSINE	25.000	0.005	60,395	0.005	0.004	40.569	695.112	73.367
C19 -----	EMULSION 7B	75.000	0.029	60,370	0.022	0.263	40.591	695.376	73.396
	* LUCITE BASE	800.000	0.059	60,295	0.136	0.280	40.727	695.656	73.455
	* EMULSION 7B	75.000	0.029	59,495	0.022	0.263	40.748	695.919	73.484
				59,420			40.748	695.919	73.484
*20TH CYCLE				59,420			40.748	695.919	73.484
				59,420			40.748	695.919	73.484
G132 --	GLASSINE	25.000	0.005	59,420	0.005	0.004	40.754	695.923	73.489
LC20	LEAD	2,000.000	2.271	59,395	1.170	35.655	41.923	731.578	75.760
G133 --	GLASSINE	25.000	0.005	57,395	0.005	0.004	41.928	731.582	75.765
				57,370			41.928	731.582	75.765
X20A	X-RAY FILM	220.000	0.043	57,370	0.032	0.392	41.960	731.973	75.808
X20B	X-RAY FILM	220.000	0.043	57,150	0.032	0.392	41.993	732.365	75.852
G134 --	GLASSINE	25.000	0.005	56,930	0.005	0.004	41.998	732.369	75.857
C20 -----	EMULSION 7B	75.000	0.029	56,905	0.022	0.263	42.019	732.632	75.886
	* LUCITE BASE	800.000	0.059	56,830	0.136	0.280	42.155	732.912	75.945

ECT MATERIALS COMPOSITION

NAME	MATERIAL	THICKNESS	MASS	HEIGHT	mfp	r.l.	Sum mfp	Sum r.l.	Sum
	NAME	µm	g/cm2	µm	%	%	%	%	g/cm2
	* EMULSION 7B	75.000	0.029	56,030	0.022	0.263	42.177	733.175	75.974
				55,955			42.177	733.175	75.974
*21ST CYCLE				55,955			42.177	733.175	75.974
				55,955			42.177	733.175	75.974
G135 -	GLASSINE	25.000	0.005	55,955	0.005	0.004	42.182	733.179	75.979
LC21	LEAD	2,000.000	2.271	55,930	1.170	35.655	43.352	768.834	78.250
G136 -	GLASSINE	25.000	0.005	53,930	0.005	0.004	43.357	768.838	78.255
				53,905			43.357	768.838	78.255
X21A	X-RAY FILM	220.000	0.043	53,905	0.032	0.392	43.389	769.229	78.298
X21B	X-RAY FILM	220.000	0.043	53,685	0.032	0.392	43.421	769.621	78.341
G137 -	GLASSINE	25.000	0.005	53,465	0.005	0.004	43.426	769.625	78.346
C21 -	EMULSION 7B	75.000	0.029	53,440	0.022	0.263	43.448	769.888	78.375
	* LUCITE BASE	800.000	0.059	53,365	0.136	0.280	43.583	770.168	78.434
	* EMULSION 7B	75.000	0.029	52,565	0.022	0.263	43.605	770.431	78.463
				52,490			43.605	770.431	78.463
*22ND CYCLE				52,490			43.605	770.431	78.463
				52,490			43.605	770.431	78.463
G138 -	GLASSINE	25.000	0.005	52,490	0.005	0.004	43.610	770.435	78.468
LC22	LEAD	2,000.000	2.271	52,465	1.170	35.655	44.780	806.090	80.739
G139 -	GLASSINE	25.000	0.005	50,465	0.005	0.004	44.785	806.094	80.744
				50,440			44.785	806.094	80.744
X22A	X-RAY FILM	220.000	0.043	50,440	0.032	0.392	44.817	806.486	80.787
X22B	X-RAY FILM	220.000	0.043	50,220	0.032	0.392	44.849	806.877	80.830
G140 -	GLASSINE	25.000	0.005	50,000	0.005	0.004	44.854	806.881	80.835
C22 -	* EMULSION 7B	75.000	0.029	49,975	0.022	0.263	44.876	807.144	80.864
	* LUCITE BASE	800.000	0.059	49,900	0.136	0.280	45.012	807.425	80.923
	* EMULSION 7B	75.000	0.029	49,100	0.022	0.263	45.033	807.688	80.952
				49,025			45.033	807.688	80.952
*23RD CYCLE				49,025			45.033	807.688	80.952
				49,025			45.033	807.688	80.952
G141 -	GLASSINE	25.000	0.005	49,025	0.005	0.004	45.038	807.692	80.957
LC23	LEAD	2,000.000	2.271	49,000	1.170	35.655	46.208	843.346	83.228
G142 -	GLASSINE	25.000	0.005	47,000	0.005	0.004	46.213	843.350	83.233
				46,975			46.213	843.350	83.233
X23A	X-RAY FILM	220.000	0.043	46,975	0.032	0.392	46.245	843.742	83.276
X23B	X-RAY FILM	220.000	0.043	46,755	0.032	0.392	46.277	844.133	83.319
G143 -	GLASSINE	25.000	0.005	46,535	0.005	0.004	46.283	844.137	83.324
C23 -	* EMULSION 7B	75.000	0.029	46,510	0.022	0.263	46.304	844.400	83.353
	* LUCITE BASE	800.000	0.059	46,435	0.136	0.280	46.440	844.681	83.412
	* EMULSION 7B	75.000	0.029	45,635	0.022	0.263	46.462	844.944	83.441
				45,560			46.462	844.944	83.441
*24TH CYCLE				45,560			46.462	844.944	83.441
				45,560			46.462	844.944	83.441
G144 -	GLASSINE	25.000	0.005	45,560	0.005	0.004	46.467	844.948	83.446
LC24	LEAD	2,000.000	2.271	45,535	1.170	35.655	47.636	880.602	85.717
G145 -	GLASSINE	25.000	0.005	43,535	0.005	0.004	47.642	880.606	85.722
				43,510			47.642	880.606	85.722
X24A	X-RAY FILM	220.000	0.043	43,510	0.032	0.392	47.674	880.998	85.765
X24B	X-RAY FILM	220.000	0.043	43,290	0.032	0.392	47.706	881.390	85.808
G146 -	GLASSINE	25.000	0.005	43,070	0.005	0.004	47.711	881.394	85.813
C24 -	* EMULSION 7B	75.000	0.029	43,045	0.022	0.263	47.732	881.657	85.842
	* LUCITE BASE	800.000	0.059	42,970	0.136	0.280	47.868	881.937	85.901
	* EMULSION 7B	75.000	0.029	42,170	0.022	0.263	47.890	882.200	85.930
				42,095			47.890	882.200	85.930
*25TH CYCLE				42,095			47.890	882.200	85.930

ECT MATERIALS COMPOSITION

NAME	MATERIAL	THICKNESS	MASS	HEIGHT	mfp	r.l.	Sum mfp	Sum r.l.	Sum
	NAME	µm	g/cm2	µm	%	%	%	%	g/cm2
				42,095			47.890	882.200	85.930
G147	GLASSINE	25.000	0.005	42,095	0.005	0.004	47.895	882.204	85.935
LC25	LEAD	2,000.000	2.271	42,070	1.170	35.655	49.065	917.859	88.206
G148	GLASSINE	25.000	0.005	40,070	0.005	0.004	49.070	917.863	88.211
				40,045			49.070	917.863	88.211
X25A	X-RAY FILM	220.000	0.043	40,045	0.032	0.392	49.102	918.254	88.254
X25B	X-RAY FILM	220.000	0.043	39,825	0.032	0.392	49.134	918.646	88.297
G149	GLASSINE	25.000	0.005	39,605	0.005	0.004	49.139	918.650	88.302
C25	* EMULSION 7B	75.000	0.029	39,580	0.022	0.263	49.161	918.913	88.331
	* LUCITE BASE	800.000	0.059	39,505	0.136	0.280	49.297	919.193	88.390
	* EMULSION 7B	75.000	0.029	38,705	0.022	0.263	49.318	919.456	88.419
G150	GLASSINE	25.000	0.005	38,630	0.005	0.004	49.323	919.460	88.424
B5(E34-E39)	CR-39	600.000	0.087	38,605	0.104	0.215	49.427	919.675	88.511
				38,005			49.427	919.675	88.511
*26TH CYCLE				38,005			49.427	919.675	88.511
				38,005			49.427	919.675	88.511
G151	GLASSINE	25.000	0.005	38,005	0.005	0.004	49.433	919.679	88.516
LC26	LEAD	2,000.000	2.271	37,980	1.170	35.655	50.602	955.333	90.787
G152	GLASSINE	25.000	0.005	35,980	0.005	0.004	50.607	955.337	90.792
				35,955			50.607	955.337	90.792
X26A	X-RAY FILM	220.000	0.043	35,955	0.032	0.392	50.639	955.729	90.835
X26B	X-RAY FILM	220.000	0.043	35,735	0.032	0.392	50.672	956.121	90.878
G153	GLASSINE	25.000	0.005	35,515	0.005	0.004	50.677	956.125	90.883
C26	* EMULSION 7B	75.000	0.029	35,490	0.022	0.263	50.698	956.388	90.912
	* LUCITE BASE	800.000	0.059	35,415	0.136	0.280	50.834	956.668	90.971
	* EMULSION 7B	75.000	0.029	34,615	0.022	0.263	50.856	956.931	91.000
				34,540			50.856	956.931	91.000
*27TH CYCLE				34,540			50.856	956.931	91.000
				34,540			50.856	956.931	91.000
G154	GLASSINE	25.000	0.005	34,540	0.005	0.004	50.861	956.935	91.005
LC27	LEAD	2,000.000	2.271	34,515	1.170	35.655	52.030	992.590	93.276
G155	GLASSINE	25.000	0.005	32,515	0.005	0.004	52.036	992.594	93.281
				32,490			52.036	992.594	93.281
X27A	X-RAY FILM	220.000	0.043	32,490	0.032	0.392	52.068	992.985	93.325
X27B	X-RAY FILM	220.000	0.043	32,270	0.032	0.392	52.100	993.377	93.368
G156	GLASSINE	25.000	0.005	32,050	0.005	0.004	52.105	993.381	93.373
C27	* EMULSION 7B	75.000	0.029	32,025	0.022	0.263	52.127	993.644	93.402
	* LUCITE BASE	800.000	0.059	31,950	0.136	0.280	52.262	993.924	93.461
	* EMULSION 7B	75.000	0.029	31,150	0.022	0.263	52.284	994.187	93.490
				31,075			52.284	994.187	93.490
*28TH CYCLE				31,075			52.284	994.187	93.490
				31,075			52.284	994.187	93.490
G157	GLASSINE	25.000	0.005	31,075	0.005	0.004	52.289	994.191	93.495
LC28	LEAD	2,000.000	2.271	31,050	1.170	35.655	53.459	1,029.846	95.766
G158	GLASSINE	25.000	0.005	29,050	0.005	0.004	53.464	1,029.850	95.771
				29,025			53.464	1,029.850	95.771
X28A	X-RAY FILM	220.000	0.043	29,025	0.032	0.392	53.496	1,030.241	95.814
X28B	X-RAY FILM	220.000	0.043	28,805	0.032	0.392	53.528	1,030.633	95.857
G159	GLASSINE	25.000	0.005	28,585	0.005	0.004	53.533	1,030.637	95.862
C28	* EMULSION 7B	75.000	0.029	28,560	0.022	0.263	53.555	1,030.900	95.891
	* LUCITE BASE	800.000	0.059	28,485	0.136	0.280	53.691	1,031.180	95.950
	* EMULSION 7B	75.000	0.029	27,685	0.022	0.263	53.712	1,031.443	95.979
				27,610			53.712	1,031.443	95.979
				27,610			53.712	1,031.443	95.979
*29TH CYCLE				27,610			53.712	1,031.443	95.979

ECT MATERIALS COMPOSITION

NAME	MATERIAL	THICKNESS	MASS	HEIGHT	mfp	r.l.	Sum mfp	Sum r.l.	Sum
	NAME	µm	g/cm2	µm	%	%	%	%	g/cm2
				27,610			53.712	1,031.443	95.979
G160	GLASSINE	25.000	0.005	27,610	0.005	0.004	53.717	1,031.447	95.984
LC29	LEAD + TLD	2,000.000	2.271	27,585	1.170	35.655	54.887	1,067.102	98.255
G161	GLASSINE	25.000	0.005	25,585	0.005	0.004	54.892	1,067.106	98.260
				25,560			54.892	1,067.106	98.260
X29A	X-RAY FILM	220.000	0.043	25,560	0.032	0.392	54.924	1,067.498	98.303
X29B	X-RAY FILM	220.000	0.043	25,340	0.032	0.392	54.956	1,067.889	98.346
G162	GLASSINE	25.000	0.005	25,120	0.005	0.004	54.962	1,067.893	98.351
C29	* EMULSION 7B	75.000	0.029	25,170	0.022	0.263	54.983	1,068.156	98.380
	* LUCITE BASE	800.000	0.059	25,095	0.136	0.280	55.119	1,068.437	98.439
	* EMULSION 7B	75.000	0.029	24,295	0.022	0.263	55.141	1,068.700	98.468
				24,220			55.141	1,068.700	98.468
				24,220			55.141	1,068.700	98.468
*30TH CYCLE				24,220			55.141	1,068.700	98.468
				24,220			55.141	1,068.700	98.468
G163	GLASSINE	25.000	0.005	24,220	0.005	0.004	55.146	1,068.704	98.473
LC30	LEAD	2,000.000	2.271	24,195	1.170	35.655	56.315	1,104.358	100.744
G164	GLASSINE	25.000	0.005	22,195	0.005	0.004	56.320	1,104.362	100.749
				22,170			56.320	1,104.362	100.749
X30A	X-RAY FILM	220.000	0.043	22,170	0.032	0.392	56.353	1,104.754	100.792
X30B	X-RAY FILM	220.000	0.043	21,950	0.032	0.392	56.385	1,105.145	100.835
G165	GLASSINE	25.000	0.005	21,730	0.005	0.004	56.390	1,105.149	100.840
C30	* EMULSION 7B	75.000	0.029	21,705	0.022	0.263	56.411	1,105.412	100.869
	* LUCITE BASE	800.000	0.059	21,630	0.136	0.280	56.547	1,105.693	100.928
	* EMULSION 7B	75.000	0.029	20,830	0.022	0.263	56.569	1,105.956	100.957
				20,755			56.569	1,105.956	100.957
				20,755			56.569	1,105.956	100.957
*31ST CYCLE				20,755			56.569	1,105.956	100.957
				20,755			56.569	1,105.956	100.957
G166	GLASSINE	25.000	0.005	20,755	0.005	0.004	56.574	1,105.960	100.962
LC31	LEAD	2,000.000	2.271	20,730	1.170	35.655	57.744	1,141.614	103.233
G167	GLASSINE	25.000	0.005	18,730	0.005	0.004	57.749	1,141.618	103.238
				18,705			57.749	1,141.618	103.238
X31A	X-RAY FILM	220.000	0.043	18,705	0.032	0.392	57.781	1,142.010	103.281
X31B	X-RAY FILM	220.000	0.043	18,485	0.032	0.392	57.813	1,142.402	103.324
G168	GLASSINE	25.000	0.005	18,265	0.005	0.004	57.818	1,142.406	103.329
C31	* EMULSION 7B	75.000	0.029	18,240	0.022	0.263	57.840	1,142.669	103.358
	* LUCITE BASE	800.000	0.059	18,165	0.136	0.280	57.976	1,142.949	103.417
	* EMULSION 7B	75.000	0.029	17,365	0.022	0.263	57.997	1,143.212	103.446
				17,290			57.997	1,143.212	103.446
				17,290			57.997	1,143.212	103.446
*32ND CYCLE				17,290			57.997	1,143.212	103.446
				17,290			57.997	1,143.212	103.446
G169	GLASSINE	25.000	0.005	17,290	0.005	0.004	58.002	1,143.216	103.451
LC32	LEAD	2,000.000	2.271	17,265	1.170	35.655	59.172	1,178.871	105.722
G170	GLASSINE	25.000	0.005	15,265	0.005	0.004	59.177	1,178.875	105.727
				15,240			59.177	1,178.875	105.727
X32A	X-RAY FILM	220.000	0.043	15,240	0.032	0.392	59.209	1,179.266	105.770
X32B	X-RAY FILM	220.000	0.043	15,020	0.032	0.392	59.241	1,179.658	105.813
G171	GLASSINE	25.000	0.005	14,800	0.005	0.004	59.246	1,179.662	105.818
C32	* EMULSION 7B	75.000	0.029	14,775	0.022	0.263	59.268	1,179.925	105.847
	* LUCITE BASE	800.000	0.059	14,700	0.136	0.280	59.404	1,180.205	105.906
	* EMULSION 7B	75.000	0.029	13,900	0.022	0.263	59.425	1,180.468	105.935
G172	GLASSINE	25.000	0.005	13,825	0.005	0.004	59.431	1,180.472	105.940
B6(E12-E05)	CR-39	600.000	0.087	13,800	0.104	0.215	59.535	1,180.687	106.027

ECT MATERIALS COMPOSITION

NAME	MATERIAL	THICKNESS	MASS	HEIGHT	mfp	r.l.	Sum mfp	Sum r.l.	Sum
	NAME	µm	g/cm2	µm	%	%	%	%	g/cm2
				13,200			59.535	1,180.687	106.027
*33RD CYCLE				13,200			59.535	1,180.687	106.027
				13,200			59.535	1,180.687	106.027
G173 -	GLASSINE	25.000	0.005	13,200	0.005	0.004	59.540	1,180.691	106.032
LC33	LEAD	2,000.000	2.271	13,175	1.170	35.655	60.709	1,216.345	108.303
G174 -	GLASSINE	25.000	0.005	11,175	0.005	0.004	60.715	1,216.349	108.308
				11,150			60.715	1,216.349	108.308
X33A	X-RAY FILM	220.000	0.043	11,150	0.032	0.392	60.747	1,216.741	108.352
X33B	X-RAY FILM	220.000	0.043	10,930	0.032	0.392	60.779	1,217.133	108.395
G175 -	GLASSINE	25.000	0.005	10,710	0.005	0.004	60.784	1,217.137	108.400
C33 -	* EMULSION 7B	75.000	0.029	10,685	0.022	0.263	60.806	1,217.400	108.429
	* LUCITE BASE	800.000	0.059	10,610	0.136	0.280	60.941	1,217.680	108.488
	* EMULSION 7B	75.000	0.029	9,810	0.022	0.263	60.963	1,217.943	108.517
				9,735			60.963	1,217.943	108.517
*34TH CYCLE				9,735			60.963	1,217.943	108.517
				9,735			60.963	1,217.943	108.517
G176 -	GLASSINE	25.000	0.005	9,735	0.005	0.004	60.968	1,217.947	108.522
LC34	LEAD + TLD	2,000.000	2.271	9,710	1.170	35.655	62.138	1,253.602	110.793
G177 -	GLASSINE	25.000	0.005	7,710	0.005	0.004	62.143	1,253.606	110.798
				7,685			62.143	1,253.606	110.798
X34A	X-RAY FILM	220.000	0.043	7,685	0.032	0.392	62.175	1,253.997	110.841
X34B	X-RAY FILM	220.000	0.043	7,465	0.032	0.392	62.207	1,254.389	110.884
G178 -	GLASSINE	25.000	0.005	7,245	0.005	0.004	62.212	1,254.393	110.889
C34 -	* EMULSION 7B	75.000	0.029	7,220	0.022	0.263	62.234	1,254.656	110.918
	* LUCITE BASE	800.000	0.059	7,145	0.136	0.280	62.370	1,254.936	110.977
	* EMULSION 7B	75.000	0.029	6,420	0.022	0.263	62.391	1,255.199	111.006
				6,345			62.391	1,255.199	111.006
*35TH CYCLE				6,345			62.391	1,255.199	111.006
				6,345			62.391	1,255.199	111.006
G179 -	GLASSINE	25.000	0.005	6,345	0.005	0.004	62.396	1,255.203	111.011
LC35	LEAD + THERM	2,000.000	2.271	6,320	1.170	35.655	63.566	1,290.858	113.282
G180 -	GLASSINE	25.000	0.005	4,320	0.005	0.004	63.571	1,290.862	113.287
				4,295			63.571	1,290.862	113.287
X35A	X-RAY FILM	220.000	0.043	4,295	0.032	0.392	63.603	1,291.253	113.330
X35B	X-RAY FILM	220.000	0.043	4,075	0.032	0.392	63.635	1,291.645	113.373
G181 -	GLASSINE	25.000	0.005	3,855	0.005	0.004	63.640	1,291.649	113.378
C35 -	* EMULSION 7B	75.000	0.029	3,830	0.022	0.263	63.662	1,291.912	113.407
	* LUCITE BASE	800.000	0.059	3,755	0.136	0.280	63.798	1,292.192	113.466
	* EMULSION 7B	75.000	0.029	2,955	0.022	0.263	63.820	1,292.455	113.495
				2,880			63.820	1,292.455	113.495
*****	*****			2,880			63.820	1,292.455	113.495
**** END OF	CALORIMETER			2,880			63.820	1,292.455	113.495
*****	*****			2,880			63.820	1,292.455	113.495
				2,880			63.820	1,292.455	113.495
G182 -	GLASSINE	25.000	0.005	2,880	0.005	0.004	63.825	1,292.459	113.500
C36 -	* EMULSION 7B	200.000	0.772	2,855	0.058	0.702	63.882	1,293.161	114.272
	* LUCITE BASE	300.000	0.035	2,655	0.051	0.105	63.933	1,293.266	114.307
	* EMULSION 7B	200.000	0.772	2,355	0.058	0.702	63.991	1,293.968	115.079
G183 -	GLASSINE	25.000	0.005	2,155	0.005	0.004	63.996	1,293.972	115.084
B7(E04-E01)	CR39	600.000	0.087	2,130	0.104	0.215	64.100	1,294.186	115.171
G184 -	GLASSINE	25.000	0.005	1,530	0.005	0.004	64.105	1,294.190	115.176
		0.000		1,505			64.105	1,294.190	115.176
*TEST ITEMS		0.000		1,505			64.105	1,294.190	115.176
		0.000		1,505			64.105	1,294.190	115.176
G185 -	GLASSINE	25.000	0.005	1,505	0.005	0.004	64.110	1,294.194	115.181

ECT MATERIALS COMPOSITION

NAME	MATERIAL	THICKNESS	MASS	HEIGHT	mfp	r.l.	Sum mfp	Sum r.l.	Sum
	NAME	μm	g/cm ²	μm	%	%	%	%	g/cm ²
S-1 —	SEALON FILM	100.000	0.015	1,480	0.011	0.035	64.121	1,294.229	115.196
X1 TEST	X-RAY FILM	220.000	0.043	1,380	0.032	0.392	64.153	1,294.621	115.239
G186 —	GLASSINE	25.000	0.005	1,160	0.005	0.004	64.158	1,294.625	115.244
S-2 —	SEALON FILM	100.000	0.015	1,135	0.011	0.035	64.169	1,294.660	115.258
X2 TEST	X-RAY FILM	220.000	0.043	1,035	0.032	0.392	64.201	1,295.051	115.301
G187 —	GLASSINE	25.000	0.005	815	0.005	0.004	64.206	1,295.055	115.306
S-3 —	SEALON FILM	100.000	0.015	790	0.011	0.035	64.217	1,295.090	115.321
X3 TEST	X-RAY FILM	220.000	0.043	690	0.032	0.392	64.249	1,295.482	115.364
G188 —	GLASSINE	25.000	0.005	470	0.005	0.004	64.254	1,295.486	115.369
S-4 —	SEALON FILM	100.000	0.015	445	0.011	0.035	64.265	1,295.521	115.384
X4 TEST	X-RAY FILM	220.000	0.043	345	0.032	0.392	64.297	1,295.913	115.427
G189 —	GLASSINE	25.000	0.005	125	0.005	0.004	64.302	1,295.916	115.432
S-5 —	SEALON FILM	100.000	0.015	100	0.011	0.035	64.313	1,295.951	115.446
		0.000	0.000	0			64.313	1,295.951	115.446
*****	*****								
BOTTOM OF THE CHAMBER									
*****	*****								